

GenCore version 5.1.7
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OM nucleic - nucleic search, using sw model

Run on: February 17, 2006, 09:49:14 ; Search time 1478 Seconds
(without alignments)
11277.670 Million cell updates/sec

Title: US-10-607-806-1-C7256_COPY_7000_9500

Perfect score: 2499

Sequence: 1 gtcgtgcacacgtcgtccag.....tttgagaccagcctcgacaa 2501

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 4996997 seqs, 3332346308 residues

Total number of hits satisfying chosen parameters: 9993994

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : N Geneseq_21:*

1: geneseqn1980s:*
2: geneseqn1990s:*
3: geneseqn2000s:*
4: geneseqn2001as:*
5: geneseqn2001bs:*
6: geneseqn2002as:*
7: geneseqn2002bs:*
8: geneseqn2003as:*
9: geneseqn2003bs:*
10: geneseqn2003cs:*
11: geneseqn2003ds:*
12: geneseqn2004as:*
13: geneseqn2004bs:*
14: geneseqn2005s:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	2498.6	100.0	12174	12 ADI35082	ADI35082 Human PLA
2	2497.4	99.9	12174	12 ADJ09983	Adj09983 Human pho
3	2497.4	99.9	12612	6 ABK47376	ABK47376 Human pho
4	456.6	18.3	42360	13 ABD33466	ABD33466 Human can
5	434.2	17.4	93544	13 ABD33504	ABD33504 Human can
6	434.2	17.4	160482	11 ACN43914	ACn43914 Human gen
7	433.2	17.3	91760	11 ACN44410	ACn44410 Human gen
8	425.4	17.0	38538	13 ABD33357	ABd33357 Human can
9	417.2	16.7	122888	6 ABK83569	ABk83569 Human can
10	416.2	16.7	6405	5 ABF97850	ABf97850 Human neu
11	416	16.6	128990	12 ADP13332	ADp13332 Renal cel
12	414.2	16.6	196686	11 ACN41170	ACn41170 Human gen
13	413.6	16.6	86000	12 ADP68568	ADp68568 Human ppa
14	413.6	16.6	215974	12 ADQ97523	ADq97523 Human can
15	413.4	16.5	13670	6 AAL42369	AAl42369 Human gua
16	413.2	16.4	6519	5 ABA15909	ABa15909 Human ner
17	408.8	16.4	17245	4 AAK83897	AAk83897 Human imm
18	406.2	16.3	348101	12 ADQ97146	ADq97146 Human can
19	405.8	16.2	227246	13 ABD33272	ABd33272 Human can

C	20	404	16.2	95240	10 ADL13556	ADl13556 Osteoarth
C	21	403.6	16.2	23639	6 ABQ78991	ABq78991 Human pho
C	22	400.4	16.0	14176	4 AAS26670	AAs26670 Human gen
C	23	400.4	16.0	14176	8 ABX74019	ABx74019 Human nov
C	24	400.4	16.0	177531	8 ACF62732	ACf62732 Cancer ba
C	25	400.4	16.0	177531	8 ADB20847	ADb20847 MRP1 base
C	26	400.4	16.0	177531	10 ADB87936	ADb87936 Human UGT
C	27	400.4	16.0	177531	10 ADB86919	ADb86919 Human MDR
C	28	400.4	16.0	177531	10 ADH92110	ADh92110 Human MDR
C	29	400.4	16.0	177531	10 ADH74617	ADh74617 Human BAC
C	30	399	16.0	87687	11 ACN45166	ACn45166 Human gen
C	31	399	16.0	131078	14 ADX06911	ADx06911 Cyclin-de
C	32	398.6	16.0	174448	11 ACN43946	ACn43946 Human gen
C	33	398.4	15.9	167343	6 ABL64403	ABl64403 Stomach c
C	34	398.4	15.9	167343	4 ABL64239	ABl64239 Thyroid c
C	35	394.8	15.8	30620	4 ABK65931	ABk65931 Human imm
C	36	394.6	15.8	19300	12 ADP74371	ADp74371 Human X c
C	37	392.4	15.7	9289	10 ADB84033	ADb84033 5' regula
C	38	391.4	15.7	226475	9 AAD58279	AAd58279 Human tum
C	39	390	15.6	13224	4 AAS41751	AAa41751 Genomic s
C	40	390	15.6	13224	4 ABA06811	ABa06811 Human gen
C	41	390	15.6	13224	6 ABV84148	ABv84148 Human pol
C	42	390	15.6	36221	4 AAS00624	AAa00624 Human dea
C	43	389.6	15.6	23456	13 ABD33110	ABd33110 Human can
C	44	389.6	15.6	60057	11 ACN44314	ACn44314 Human can
C	45	388	15.5	75252	11 ACN44450	ACn44450 Human gen

ALIGNMENTS

RESULT 1	
ID ADI35082	ADI35082 standard; DNA; 12174 BP.
XX	
AC	ADI35082;
XX	
DT	22-APR-2004 (first entry)
XX	
DE	Human PLA2G1B nucleotide sequence.
XX	
KW	PLA2G1B ; fat deposition; leanness; polymorphism;
KW	non-insulin dependent diabetes mellitus; NIDDM; hyperinsulinemia;
KW	hyperextension; glucose intolerance; dyslipidemia; hypercoagulability;
KW	microalbuminuria; human; gene; ds.
XX	
OS	Homo sapiens.
XX	
PN	W02004002295-A2.
XX	
PD	08-JAN-2004.
XX	
PE	27-JUN-2003; 2003WO-US020830.
XX	
PR	27-JUN-2002; 2002US-0392361P.
XX	
PA	(SEQU-) SEQUENOM INC.
XX	
PI	Adam GIR, Langdown ML;
XX	
DR	WPI; 2004-082843/08.
XX	
PT	P-PSDB; ADI35083.
XX	
PS	Diagnosing a predisposition to fat deposition or leanness, useful for
XX	diagnosing a predisposition to e.g. diabetes or hypertension, comprises
CC	detecting the presence of a polymorphism in the PLA2G1B nucleic acid from
CC	the subject.
CC	Claim 1; SEQ ID NO 1; 91pp: English.
CC	The invention relates to diagnosing a predisposition to fat deposition or
CC	leanness in a subject comprising detecting the presence or absence of a
CC	polymorphic variation associated with fat deposition at a polymorphic

CC site in a PLA2G1B nucleotide sequence in a nucleic acid sample from a
CC subject, where the presence of the polymorphic variation indicates a
CC predisposition to fat deposition in the subject. The polymorphic
CC variation is a guanine at position 7328 or thymine at position 9182 of
CC the present sequence. The method is useful for diagnosing a
CC predisposition to fat deposition or leanness in a subject, and
CC consequently for diagnosing a predisposition to non-insulin dependent
CC diabetes mellitus (NIDDM) in a subject and conditions such as
CC hyperinsulinemia, hypertension, glucose intolerance, dyslipidemia,
CC hypercoagulability, or microalbuminuria, which can lead to early
CC prescription of preventive measures. The present sequence represents a
CC human PLA2G1B nucleotide sequence.

XX Sequence 12174 BP; 3217 A; 2992 C; 2738 G; 3215 T; 0 U; 12 Other;

Query Match 100.0%; Score 2499; DB 12; Length 12174;

Best Local Similarity 99.9%; Pred. No. 0;

Matches 2500; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

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QY 1 GTCTGTGACAGCTGCTCCAGCCTGGGTAAACAGACAACTCTGTCTCAAAAAAATG 60
Db 7000 GTCTGTGACAGCTGCTCCAGCCTGGGTAAACAGACAACTCTGTCTCAAAAAAATG 7059
QY 61 CTTTCAATTAATATATGATTAATAAGAGCTTAATTTTCAAGCCATAGGATATTCTCC 120
Db 7060 CTTTCAATTAATATATGATTAATAAGAGCTTAATTTTCAAGCCATAGGATATTCTCC 7119
QY 121 TGAAGCATCTTGCGCAAGTCATCCCACTGTCTCTGAGAGTGCGCAGGTGAGC 180
Db 7120 TGAAGCATCTTGCGCAAGTCATCCCACTGTCTCTGAGAGTGCGCAGGTGAGCCTGAC 7179
QY 181 CTATTGCTCTGACATTAATCTCCATCTCAGCTGTCCCTCCACTTTCCAGGTGCTGCCGA 240
Db 7180 CTATTGCTCTGACATTAATCTCCATCTCAGCTGTCCCTCCACTTTCCAGGTGCTGCCGA 7239
QY 241 CACATGACAACTGTACGACGACGACGACGACGACGACGACGACGACGACGACGACGAC 300
Db 7240 CACATGACAACTGTACGACGACGACGACGACGACGACGACGACGACGACGACGACGAC 7299
QY 301 MMCCGTACACCCACA CTAATTCATACTCTGCTGTGCGCTGCGCAATCACTGTAGCACTA 360
Db 7300 MMCCGTACACCCACA CTAATTCATACTCTGCTGTGCGCTGCGCAATCACTGTAGCACTA 7359
QY 361 GGTATTATCCTTCTTGAAGCTATGAAATTTAGTGGTTCTCAGTAGCCGGGGGGAATA 420
Db 7360 GGTATTATCCTTCTTGAAGCTATGAAATTTAGTGGTTCTCAGTAGCCGGGGGGAATA 7419
QY 421 ATAGTAACAAGCAGCATGATTAAGTAAATTTCTTGGTCTGCGCACTGTCTCTTTA 480
Db 7420 ATAGTAACAAGCAGCATGATTAAGTAAATTTCTTGGTCTGCGCACTGTCTCTTTA 7479
QY 481 ATCTCTAGAAACA CTAATGGAATAGTAACAATTAATCTCACTTAACAGATAAAGAACT 540
Db 7480 ATCTCTAGAAACA CTAATGGAATAGTAACAATTAATCTCACTTAACAGATAAAGAACT 7539
QY 541 GAGGCTCAGAAAGCTGAGCTATTTGGCCAAAGTACACAGCTTTGTAAGTGGTGAAGTT 600
Db 7540 GAGGCTCAGAAAGCTGAGCTATTTGGCCAAAGTACACAGCTTTGTAAGTGGTGAAGTT 7599
QY 601 GGGTTTTTTTTTGTGTGTTTGAAGACAGGGCTTGTCTGTCAACCCAGGATGAGAC 660
Db 7600 GGGTTTTTTTTTGTGTGTTTGAAGACAGGGCTTGTCTGTCAACCCAGGATGAGAC 7659
QY 661 AGTGTGCAACCAATAGTCACTGCAAGCTCAACCTCTGAGCTCAAGGAGTCTGTGACC 720
Db 7660 AGTGTGCAACCAATAGTCACTGCAAGCTCAACCTCTGAGCTCAAGGAGTCTGTGACC 7719
QY 721 TGAAGCTCCCAAGTACGTGGGAACTAGAGGTGACACACAGCCTGTGGCTAATTAAGAA 780
Db 7720 TGAAGCTCCCAAGTACGTGGGAACTAGAGGTGACACACAGCCTGTGGCTAATTAAGAA 7779
QY 781 ATTTTGTGAAGACATGGGCTTACTAGCTTGGCCAGGCTTCTTAAACTCTGGCTT 840
Db 781 ATTTTGTGAAGACATGGGCTTACTAGCTTGGCCAGGCTTCTTAAACTCTGGCTT 840
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Db 7780 ATTTTGTGAAGACATGGGCTTACTAGCTTGGCCAGGCTTCTTAAACTCTGGCTT 7839
QY 841 CAAGCAATCCCTCCATCTGGGCAATCCCAAGTGTGGGATTTACAGGGGTGAGCACCATG 900
Db 7840 CAAGCAATCCCTCCATCTGGGCAATCCCAAGTGTGGGATTTACAGGGGTGAGCACCATG 7899
QY 901 TGGCGTACTATTATTTCTTACATTCATCTTCCAAATGAATGAATCCAGAAACAG 960
Db 7900 TGGCGTACTATTATTTCTTACATTCATCTTCCAAATGAATGAATCCAGAAACAG 7959
QY 961 GGATTACTGCTATTTTCTTCTTCTTTTGAAGACAGATCTCACTTATCACTCA 1020
Db 7960 GGATTACTGCTATTTTCTTCTTCTTTTGAAGACAGATCTCACTTATCACTCA 8019
QY 1021 CTTCCGTGAGTACCTGGAATTAACAGCTGACACACATGCTTGGCTAATTTTGTATTT 1140
Db 8080 CTTCCGTGAGTACCTGGAATTAACAGCTGACACACATGCTTGGCTAATTTTGTATTT 8139
QY 1141 TAGCAGAGATGGGTTTACCATGTTGCCAGGCTGCTCAAACTCTGACCTCAAGTG 1200
Db 8140 TAGCAGAGATGGGTTTACCATGTTGCCAGGCTGCTCAAACTCTGACCTCAAGTG 8199
QY 1201 ATCTGCGCTCCAGTCCCAAGTGTGGAATTAATGAGGAGTCACTGTGCTGGC 1260
Db 8200 ATCTGCGCTCCAGTCCCAAGTGTGGAATTAATGAGGAGTCACTGTGCTGGC 8259
QY 1261 CGATTACTGCTATTTTCTTATTTGCTATATCCCAAGTCTAGAGCAGTGTCTGACATAT 1320
Db 8260 CGATTACTGCTATTTTCTTATTTGCTATATCCCAAGTCTAGAGCAGTGTCTGACATAT 8319
QY 1321 AGTAGTCTCAATTAATTAATGAATGAATGACAGCTGATTAATCTTTCTTTCTT 1380
Db 8320 AGTAGTCTCAATTAATTAATGAATGAATGACAGCTGATTAATCTTTCTTTCTT 8379
QY 1381 TTTTAAACAATCTTGCAACTTTGAGAAATTAATTAATTAATTAATTAATTAATTAAT 1440
Db 8380 TTTTAAACAATCTTGCAACTTTGAGAAATTAATTAATTAATTAATTAATTAATTAAT 8439
QY 1441 CTTATCACCTGTTATGACCTTTTCAATATGCTCAAACTTTATTTGTTACTGTTTTTC 1500
Db 8440 CTTATCACCTGTTATGACCTTTTCAATATGCTCAAACTTTATTTGTTACTGTTTTTC 8499
QY 1501 ATTTGTTACTATTTATGCACTGAATTAATGCTTAATTTGTTATATCACTCTGCTC 1560
Db 8500 ATTTGTTACTATTTATGCACTGAATTAATGCTTAATTTGTTATATCACTCTGCTC 8559
QY 1561 CACTTTAAGGCCAAATTTCAATCTGATGAAGCTATGAACCTCTCCCAAGAAA 1620
Db 8560 CACTTTAAGGCCAAATTTCAATCTGATGAAGCTATGAACCTCTCCCAAGAAA 8619
QY 1621 TACACACACACACACACTCAACAGTTTTTTTTTAACTTTTGAACCTAAGACAAG 1680
Db 8620 TACACACACACACACACTCAACAGTTTTTTTTTAACTTTTGAACCTAAGACAAG 8679
QY 1681 AACCTGATTAAGAGATGTTTGTCAATTAATTAATTAATTAATTAATTAATTAAT 1740
Db 8680 AACCTGATTAAGAGATGTTTGTCAATTAATTAATTAATTAATTAATTAATTAAT 8739
QY 1741 CTCAAGCCTGTAACCAAGTACTTTGGAAGTCAAGGTGGGAGTCACTTAGGTTGAGA 1800
Db 8740 CTCAAGCCTGTAACCAAGTACTTTGGAAGTCAAGGTGGGAGTCACTTAGGTTGAGA 8799
QY 1801 AGTTGAGACCAAGCTGTCAATATGTTGAACCTTATCTTAATTAATTAATTAATTAAT 1860
Db 8800 AGTTGAGACCAAGCTGTCAATATGTTGAACCTTATCTTAATTAATTAATTAATTAAT 8859
QY 1861 AGCTGGGTGTAGTATGATGATGCTGTAGTCCAGCTACTCGGAGGCTGAGGCAGAGAA 1920
Db 8860 AGCTGGGTGTAGTATGATGATGCTGTAGTCCAGCTACTCGGAGGCTGAGGCAGAGAA 8919
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QY 1921 TTGCTGAACTGGAGGAGAGGTTGAGGCGAGATCCCACTGCACTCCAGC 1980
DB 8920 TTGCTTGAACCTGGAGGAGAGGTTGAGGCGAGATCCCACTGCACTCCAGC 8979
QY 1981 CTGGGCGGACAGAGGAGAGCTCTATCTCAAAAAAATAAATAAATAAAGATCGAG 2040
DB 8980 CTGGGCGGACAGAGGAGAGCTCTATCTCAAAAAAATAAATAAATAAAGATCGAG 9039
QY 2041 AGAAACAAACTAATTAAGATTCTGAAGTAAAGAGATACGTAAATTATATGTAATTA 2100
DB 9040 AGAAACAAACTAATTAAGATTCTGAAGTAAAGAGATACGTAAATTATATGTAATTA 9099
QY 2101 AGTTTAAATGCAATTTAATCTGAATCTTAATGTTTATTTGTTTAAAGTAAAGCAAGC 2160
DB 9100 AGTTTAAATGCAATTTAATCTGAATCTTAATGTTTATTTGTTTAAAGTAAAGCAAGC 9159
QY 2161 CAAAAGTAATGCAATTTAATCTGAATCTTAATGTTTATTTGTTTAAAGTAAAGCAAGC 2220
DB 9160 CAAAAGTAATGCAATTTAATCTGAATCTTAATGTTTATTTGTTTAAAGTAAAGCAAGC 9219
QY 2221 ATATCTCTACTAATCCCAAGATTAACAGTATATCTTCCAGATTTTGGGGCATACAC 2280
DB 9220 ATATCTCTACTAATCCCAAGATTAACAGTATATCTTCCAGATTTTGGGGCATACAC 9279
QY 2281 TAGCTTTTATTTGGGAAATTTTCAATGTCAGGAGCATACCTAATTTTCTAAATGTC 2340
DB 9280 TAGCTTTTATTTGGGAAATTTTCAATGTCAGGAGCATACCTAATTTTCTAAATGTC 9339
QY 2341 ATGTAGTATTCATTTAAGATGTTCCATTAATTTTAAATAATACATGCTTTAAAGTAGA 2400
DB 9340 ATGTAGTATTCATTTAAGATGTTCCATTAATTTTAAATAATACATGCTTTAAAGTAGA 9399
QY 2401 AACTAGGTTGGGCAATGTCGCTCAAGCTGTATCCAGCATTTGGAGGCCAGGCAAA 2460
DB 9400 AACTAGGTTGGGCAATGTCGCTCAAGCTGTATCCAGCATTTGGAGGCCAGGCAAA 9459
QY 2461 TGGATCACTTGGAGGCCAGGATTTGAGAGCGAGCTGAGCAA 2501
DB 9460 TGGATCACTTGGAGGCCAGGATTTGAGAGCGAGCTGAGCAA 9500

RESULT 2
ADJ09983 ID ADJ09983 standard; DNA; 12174 BP.
AC ADJ09983;
XX 17-JUN-2004 (first entry)
XX Human phospholipase A2 (PLA2G1B) DNA SeqID 1.
DE
XX human; gene; ds; fat reduction; fat deposition; phospholipase A2;
KM PLA2G1B; chromosome 12q24; single nucleotide polymorphism; SNP;
KM appetite suppressant; lipase inhibitor; exercise regimen; obesity;
KM non-insulin dependent diabetes mellitus; NIDDM; cardiovascular disorder;
KM hypertension; antidiabetic.
XX Homo sapiens.
OS
XX Key Location/Qualifiers
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FT variation
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FT variation
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PN WO2004002296-A2.
XX 08-JAN-2004.
PD 27-JUN-2003; 2003WO-US020831.
XX 27-JUN-2002; 2002US-0392362P.
PR 27-JUN-2002; 2002US-0392362P.
XX (SEQID-) SEQUENCE INC.
PA Adam GIR, Langdown ML, Denisenko MF, Dennis E, Cantor C;
PI Rubin B;
XX WPI, 2004-071944/07.
DR P-PSDB; ADJ09984.
XX Identifying a candidate therapeutic for fat reduction, useful for
PT treating diabetes, by introducing a test molecule to a system comprising
PT PLA2G1B protein or nucleic acid, and determining the presence of
PT interaction between the compounds.
XX Claim 1; SEQ ID NO 1; 116pp; English.
PS This invention relates to a novel candidate therapeutic agent useful for
XX fat reduction and disorders related to fat depositions. Specifically, it
XX refers to polymorphic variations in the phospholipase A2 (PLA2G1B) DNA,
XX which is located on chromosome 12q24 and has been associated with central
XX fat deposition. The present invention describes methods to detect the
XX presence or absence of these single nucleotide polymorphisms of PLA2G1B,
XX in particular G7328A and T9182G, and subsequently provide treatment that
XX reduces fat deposition. This treatment may consist of an appetite
XX suppressant, a lipase inhibitor, a phospholipase inhibitor, an exercise
XX regimen, a dietary regimen, psychological counseling, psychotherapy or a
XX psychotherapeutic. Accordingly, PLA2G1B is a target for reducing fat
XX deposition and it can be used to treat both obesity and non-insulin
XX dependent diabetes mellitus (NIDDM), as well as cardiovascular disorders
XX such as hypertension. As such, it exhibits antidiabetic activity. This
XX polynucleotide sequence is the human PLA2G1B DNA of the invention.
SQ Sequence 12174 BP; 3220 A; 2996 C; 2739 G; 3219 T; 0 U; 0 Other;
Query Match 99.9%; Score 2497.4; DB 12; Length 12174;
```

Best Local Similarity 99.8%; Pred. No. 0;
Matches 2495; Conservative 5; Mismatches 1; Indels 0; Gaps 0;

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Qy 1 GTCGTGCTACGCTGCTGCAAGCTTGGTAAAGAGCACTGTGTCTCAAAAAAATG 60
Db 7000 GTCCGTGCTACGCTGCTGCAAGCTTGGTAAAGAGCACTGTGTCTCAAAAAAATG 7059
Qy 61 CTTTCATTAATATATATATATATATATATATATATATATATATATATATATAT 120
Db 7060 CTTTCATTAATATATATATATATATATATATATATATATATATATATATATAT 7119
Qy 121 TGAAGACTTGGGAGAGTCAATCCCACTGTCTCTGAGAGTGGCAGGTGAAGGCTGAC 180
Db 7120 TGAAGACTTGGGAGAGTCAATCCCACTGTCTCTGAGAGTGGCAGGTGAAGGCTGAC 7179
Qy 181 CTATGTCTGTGCACTTACTCTATCTGAGCTGTCCCTCCACTTTCAGAGTGTCCAGA 240
Db 7180 CTATGTCTGTGCACTTACTCTATCTGAGCTGTCCCTCCACTTTCAGAGTGTCCAGA 7239
Qy 241 CACATGAACCTGTGAGACCAAGGCAAGAGCTGGAAGCTGTAAATTTTGTGTGACA 300
Db 7240 CACATGAACCTGTGAGACCAAGGCAAGAGCTGGAAGCTGTAAATTTTGTGTGACA 7299
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Qy 361 GGTATATCCCTTCTGTTGACCTATGAAATTCATGATGCTTCTCAGTGGCCGGGGGAAATA 420
Db 7360 GGTATATCCCTTCTGTTGACCTATGAAATTCATGATGCTTCTCAGTGGCCGGGGGAAATA 7419
Qy 421 ATAGTAACAAGCAGCATATTTAGTGTAAATTTTCTGGGTCTGGGAGTGTCTCTTTA 480
Db 7420 ATAGTAACAAGCAGCATATTTAGTGTAAATTTTCTGGGTCTGGGAGTGTCTCTTTA 7479
Qy 481 ATCCCTCAGAACCACTATGAGATAGTACATTAATCTCACTTAAACAGATTAAGAAACT 540
Db 7480 ATCCCTCAGAACCACTATGAGATAGTACATTAATCTCACTTAAACAGATTAAGAAACT 7539
Qy 541 GAGGCTCAGAGGCTGAGCTATTTGGCCAAAGTCAACAGCTTGTATAGTGTGACAGTTT 600
Db 7540 GAGGCTCAGAGGCTGAGCTATTTGGCCAAAGTCAACAGCTTGTATAGTGTGACAGTTT 7599
Qy 601 GGGTTTTTTTTTGTGTGTATAGAGACAGGGGCTTGTCTGTGACCCGAGATGAGCAC 660
Db 7600 GGGTTTTTTTTTGTGTGTATAGAGACAGGGGCTTGTCTGTGACCCGAGATGAGCAC 7659
Qy 661 AGTGTGCAACCATAGTCACTGACGCTCAACCTCTCTGAGCTCAAGGATCTGTGACC 720
Db 7660 AGTGTGCAACCATAGTCACTGACGCTCAACCTCTCTGAGCTCAAGGATCTGTGACC 7719
Qy 721 TCAAGCTCCCAAGTATGCTGGGACCTAGAGGCTGACACCAAGGCTGTATATTAATAA 780
Db 7720 TCAAGCTCCCAAGTATGCTGGGACCTAGAGGCTGACACCAAGGCTGTATATTAATAA 7779
Qy 781 ATTTTGTATAGAGCTGGGTCTTACTAGTGTGGCCAGGCTGTCTTAAACTCTGGCTT 840
Db 7780 ATTTTGTATAGAGCTGGGTCTTACTAGTGTGGCCAGGCTGTCTTAAACTCTGGCTT 7839
Qy 841 CAAGCAATCTCTACCTTGGGATCCCAAGTGTGGGATTAAGGGGTGAGCCACCATG 900
Db 7840 CAAGCAATCTCTACCTTGGGATCCCAAGTGTGGGATTAAGGGGTGAGCCACCATG 7899
Qy 901 TCGGGCTACTATTTCTTACATTCATCTTTTCAATAGATATAGATTCACAGAACG 960
Db 7900 TCGGGCTACTATTTCTTACATTCATCTTTTCAATAGATATAGATTCACAGAACG 7959
Qy 961 GGAATTAATGCTATTTTCTTCTTTTGTATAGAGCAGAGTCTCATCACTCA 1020
Db 7960 GGAATTAATGCTATTTTCTTCTTTTGTATAGAGCAGAGTCTCATCACTCA 8019
Qy 1021 CCTCGTGAAGCTCACTGCAACCTGTGCTCCGGGTTCAAGATTTCTCTGCTTAAC 1080
Db 1021 CCTCGTGAAGCTCACTGCAACCTGTGCTCCGGGTTCAAGATTTCTCTGCTTAAC 1080
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Db 8020 CCTCGTGAAGCTCACTGCAACCTGTGCTCCGGGTTCAAGCATTTCTCTGCTTAAC 8079
Qy 1081 CTTCTGAGTATGCTGAATTAACAAGTGCACCACTATGCTGTGTAATTTTGTATTTT 1140
Db 8080 CTTCTGAGTATGCTGAATTAACAAGTGCACCACTATGCTGTGTAATTTTGTATTTT 8139
Qy 1141 TAGCAGAGATGGGGTTTATCCATGTGGCCAGGCTGGCTCAAACTCTGACCTCAAGTG 1200
Db 8140 TAGCAGAGATGGGGTTTATCCATGTGGCCAGGCTGGCTCAAACTCTGACCTCAAGTG 8199
Qy 1201 ATTCGCTGCTGAGTCTCCCAAGTGTGGAATTAATAGGGGTATGACATGCTGGC 1260
Db 8200 ATTCGCTGCTGAGTCTCCCAAGTGTGGAATTAATAGGGGTATGACATGCTGGC 8259
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Qy 1321 AGTAGTGTCTCAATTAATTAATGATGAATGCAAGCTTAGATATAACTTTCTTTTCTT 1380
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Qy 1381 TTTTAAAAACAATCTTGACACTTTTGCAATTTGCAATTAATCAATCTGCTTTTCA 1440
Db 8380 TTTTAAAAACAATCTTGCAACTTTTGCAATTTGCAATTAATCAATCTGCTTTTCA 8439
Qy 1441 CTATACCTTGTATGACTTTTCAATATGCTCAAACTTTATTTGTTACTGTGTTTTTC 1500
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Qy 1501 ATTTGTACTATTTTATGCTAGTAATATATGCTTTAATTTGCTTATACATCTCTGCTC 1560
Db 8500 ATTTGTACTATTTTATGCTAGTAATATATGCTTTAATTTGCTTATACATCTCTGCTC 8559
Qy 1561 CACTTTAAGAGCCCAATTTTCAATCTGATGAAGAACTATGAAACCTCTCCCAAGAAA 1620
Db 8560 CACTTTAAGAGCCCAATTTTCAATCTGATGAAGAACTATGAAACCTCTCCCAAGAAA 8619
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Db 8680 AACCTGCAATGAGAGATTTTGTTCATATTAATTAATAATACTAGTTGGCAGATGA 8739
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Db 8740 CTCAGGCTGTATACCAAGTACTTTTGAAGTCCAGGTGGGTGATCATCTTGAAGTGA 8799
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Db 8800 AGTTGAGACCAAGCTGTGCAATATGATGAACCTATCTCTAATAATAATAATAAT 8859
Qy 1861 AGCTGGGTGTATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 1920
Db 8860 AGCTGGGTGTATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 8919
Qy 1921 TTGCTTGAACCTGAGAGGAGGTTTGAAGTGAAGGCTGAGATCCCACTGACCTCCAGC 1980
Db 8920 TTGCTTGAACCTGAGAGGAGGTTTGAAGTGAAGGCTGAGATCCCACTGACCTCCAGC 8979
Qy 1981 CTGGGCGACACAGCGAGCTCTATCTCAAAAAAATTAATTAATAATAAGATCGAG 2040
Db 8980 CTGGGCGACACAGCGAGCTCTATCTCAAAAAAATTAATTAATAATAAGATCGAG 9039
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Db 9040 AGAAAACAAACTAATAAGATTCCTGAAGTGAAGAGATAGTAATTAATTAATAATA 9099
Qy 2101 AGTTTAATGCAATTTTAATGTAATCTTATTTGTTATTTTGTATTAATAAGTAACAGC 2160
Db 9100 AGTTTAATGCAATTTTAATGTAATCTTATTTGTTATTTTGTATTAATAAGTAACAGC 9159
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 Db 9280 TAGCTTTTATTTTGGGAAAATTTCCATGTCGAGGCACTACTAATTTTCTTAATGCT 9339
 QY 2341 ATGATGATTTCCATTTTAAGATGTTCCATTAATTTTAAATATACATGCTTTAAAGTAGAGA 2400
 Db 9340 ATGATGATTTCCATTTTAAGATGTTCCATTAATTTTAAATATACATGCTTTAAAGTAGAGA 9399
 QY 2401 AACTAGGTTGGGCAATGTTGGCTCAAGCCTGTATCCAGCACTTTGGAGGCCGAGCAAA 2460
 Db 9400 AACTAGGTTGGGCAATGTTGGCTCAAGCCTGTATCCAGCACTTTGGAGGCCGAGCAAA 9459
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 Db 9460 TGGATCACTTGAAGTCCGGAATTTGAGACCAAGCCTGGACAA 9500
 RESULT 3
 ABR47376
 ID ABR47376 standard; DNA; 13612 BP.
 XX ABR47376;
 AC
 XX 18-JUN-2002 (first entry)
 DT
 XX Human Phospholipase A2, groupIB (PLA2G1B) gene.
 DE
 XX Human; ds; gene; SNP; single nucleotide polymorphism; pancreatitis;
 KM pancreatic cancer; Phospholipase A2 groupIB; PLA2G1B; gene therapy;
 KW haplotype; genotype; chromosome 12q23-q24.1; transgenic; drug screening.
 XX
 OS Homo sapiens.
 XX
 FH Key Location/Qualifiers
 FT variation replace(3845,A)
 FT /*tag= a
 FT /label= SNP
 FT /note= "Single nucleotide polymorphism"
 FT variation replace(3968,A)
 FT /*tag= b
 FT /label= SNP
 FT /note= "Single nucleotide polymorphism"
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 FT /*tag= c
 FT /product= "PLA2G1B"
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 FT intron 4087..5785
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 FT /number= 9
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 FT /*tag= h
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 FT /note= "Single nucleotide polymorphism"
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 FT /number= 3

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 FT /*tag= m
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 EN W0200212562-A2.
 XX
 PD 14-FEB-2002.
 XX
 PF 06-AUG-2001; 2001WO-US024663.
 XX
 PR 04-AUG-2000; 2000US-0223179P.
 XX
 PA (GENA-) GENAISSANCE PHARM INC.
 XX
 PI Kazemi A, Kliehm SE, Koshy B;
 XX WPI; 2002-303982/34.
 DR P-PSDB; AAU78667.
 XX
 PT Novel isolated human Phospholipase A2, Group IB pancreas polynucleotide,
 PT for therapeutic purposes, for studying expression and function of the
 PT polynucleotide and for expressing the phospholipase protein.
 XX
 PS Claim 1; Fig 1; 51pp; English.
 XX
 CC The invention relates to an isolated human Phospholipase A2, Group IB
 CC (pancreas) (PLA2G1B) polynucleotide comprising a sequence which is a
 CC polymorphic variant for a reference sequence for the PLA2G1B gene or its
 CC fragment, or a polymorphic variant of a reference sequence for a PLA2G1B
 CC cDNA or its fragment. Also included are haplotyping/genotyping the
 CC PLA2G1B gene of an individual, predicting the haplotype pair for the
 CC PLA2G1B gene of an individual, identifying an association between a trait
 CC and at least one haplotype or haplotype pair of the PLA2G1B gene, an
 CC isolated genotyping oligonucleotide for detecting a polymorphism in the
 CC PLA2G1B gene, a recombinant non-human organism transformed or transfected
 CC with the PLA2G1B sequence, where the organism expresses a PLA2G1B protein
 CC encoded by the first nucleotide sequence or by the polymorphic variant
 CC sequence, an isolated polypeptide comprising a sequence which is a
 CC polymorphic variant of a reference sequence for the PLA2G1B protein or
 CC its fragment, an anti-PLA2G1B monoclonal antibody, screening for drugs
 CC targeting PLA2G1B, a computer system for storing and analyzing
 CC polymorphism data for the PLA2G1B gene and a genome anthology for PLA2G1B
 CC gene. The PLA2G1B variant is useful in studying the expression and
 CC function of PLA2G1B, and in expressing PLA2G1B protein for use in
 CC screening for candidate drugs to treat diseases related to PLA2G1B
 CC activity (e.g. pancreatitis and pancreatic cancer) and for therapeutic
 CC purposes. The transgenic organism is useful for studying expression of
 CC the PLA2G1B isogenes in vivo, for in vivo screening and testing of drugs
 CC targeted against PLA2G1B protein, and for testing the efficacy of
 CC therapeutic agents and compounds in a biological system. The antibody is
 CC useful for studying the effect of the variation on the biological
 CC activity of PLA2G1B as well as on the binding affinity of candidate drugs
 CC targeting PLA2G1B. The present sequence is the PLA2G1B gene which is
 CC located on chromosome 12q23-q24.1
 CC
 SQ Sequence 13612 BP; 3637 A; 3290 C; 3070 G; 3615 T; 0 U; 0 Other;
 Query Match 99.8%; Score 2497.4; DB 6; Length 13612;
 Best Local Similarity 99.8%; Pred. No. 0;
 Matches 2495; Conservative 5; Mismatches 1; Indels 0; Gaps 0;
 QY 1 GTCGTCTACTGCTGTCTCAGCTGGTAAACAGACAACTCTGTCCTCAAAAAAATATG 60

Db	6516	GTGCGTCACACGCTGTCACAGCGACTGCTGTCACAAAAAATG	6575
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Db	6576	CTTTCATTAATATATGATAAAGGACTTAATTTTCAAGCCATAGATCATTTCTCC	6635
Oy	121	TGAAGCATCTTGGCGAAGTCATCCCGACTGTTCTGAGTGGCGAGTGAAGGCTGAC	180
Db	6636	TGAAGCATCTTGGCGAAGTCATCCCGACTGTTCTGAGTGGCGAGTGAAGGCTGAC	6695
Oy	181	CTATTGCTCTGCACTTACTCTCATCTGACGTCTCCCTCCACTTTCAGGTGTGGCAGA	240
Db	6696	CTATTGCTCTGCACTTACTCTCATCTGACGTCTCCCTCCACTTTCAGGTGTGGCAGA	6755
Oy	241	CACATGACAACTGCTACGACACGAGCCAAAGACTGACAGCTGTAAATTTCTGTGACA	300
Db	6756	CACATGACAACTGCTACGACACGAGCCAAAGACTGACAGCTGTAAATTTCTGTGACA	6815
Oy	301	MMCCGTACACCCACACTATTCATACGTGTCTGTGGCTGGCAATCACCTGTACAGTA	360
Db	6816	AACGGTACACCCACACTATTCATACGTGTCTGTGGCTGGCAATCACCTGTACAGTA	6875
Oy	361	GGTTATCCCTTCTTGAACCTATGAATTCATGTGGTCTCAGTATGGCCGGGGGAATA	420
Db	6876	GGTTATCCCTTCTTGAACCTATGAATTCATGTGGTCTCAGTATGGCCGGGGGAATA	6935
Oy	421	ATAGTAACAAGCCATGATTTAGATTAATTTCTGGGTCGGGAGTGTCTCTTTA	480
Db	6936	ATAGTAACAAGCCATGATTTAGATTAATTTCTGGGTCGGGAGTGTCTCTTTA	6995
Oy	481	ATCTCAGAACCAACAATAATGGATAGTACAAATTAATCTCACTTAACAGATTAAGAACT	540
Db	6996	ATCTCAGAACCAACAATAATGGATAGTACAAATTAATCTCACTTAACAGATTAAGAACT	7055
Oy	541	GAGGCTCAGAAAGGCTGAGCTATTTGGCCAAAGTACACAGCTTGTAAAGTGTGACATT	600
Db	7056	GAGGCTCAGAAAGGCTGAGCTATTTGGCCAAAGTACACAGCTTGTAAAGTGTGACATT	7115
Oy	601	GGGTTTTTTTTGTGTTTGAAGAAGGGCTGTGCTGTGACCCAGGATGAGAC	660
Db	7116	GGGTTTTTTTTGTGTTTGAAGAAGGGCTGTGCTGTGACCCAGGATGAGAC	7175
Oy	661	AGTGTGCAACCATAGGTCACTGACGCTCAACTCTCTAGCTCAAGGATCTGTGACC	720
Db	7176	AGTGTGCAACCATAGGTCACTGACGCTCAACTCTCTAGCTCAAGGATCTGTGACC	7235
Oy	721	TCAGCTCCCAAGTACTGGACCTACAGCGTGGACACACAGCTGGCTAATTAAAAA	780
Db	7236	TCAGCTCCCAAGTACTGGACCTACAGCGTGGACACACAGCTGGCTAATTAAAAA	7295
Oy	781	ATTTTTTTTGAAGACTGGGTCTTAAGTGGAGGAGGCTGTCTTAAACTCTGGCTT	840
Db	7296	ATTTTTTTTGAAGACTGGGTCTTAAGTGGAGGAGGCTGTCTTAAACTCTGGCTT	7355
Oy	841	CAGCAATCTCTTCACTTGGATCCCAAGTGTGAGATTAAGGGGTGAGCCACATG	900
Db	7356	CAGCAATCTCTTCACTTGGATCCCAAGTGTGAGATTAAGGGGTGAGCCACATG	7415
Oy	901	TGGGCTACTTATTTCTTTACATTCATCTTTCAAATGAATGAATTCACAGACAG	960
Db	7416	TGGGCTACTTATTTCTTTACATTCATCTTTCAAATGAATGAATTCACAGACAG	7475
Oy	961	GGAATTAATGCTATTTCTTCTCTTTTGTGAGACAGTGTCACTTCAACCTGAA	1020
Db	7476	GGAATTAATGCTATTTCTTCTCTTTTGTGAGACAGTGTCACTTCAACCTGAA	7535
Oy	1021	CCATCGTACGCTACATGCAACTCTGACCTCCGGGTTCAAGATTTCTCTGCTAAGC	1080
Db	7536	CCATCGTACGCTACATGCAACTCTGACCTCCGGGTTCAAGATTTCTCTGCTAAGC	7595
Oy	1081	CTCTCAGTACGCTGAATTAACAAGCTGCAACAATGCTTGGCTAATTTTGTATTTT	1144

Db	7596	CTCCTGAGTAGCTGGAAATTACAAAGCGGACACAAACGTTGGCTAATTTTTGTATTTT	7655
Qy	1141	TAGCAGAGATGGGGTTTTTACATATTTGCCAGAGGCTGGTCTCAAACTCTCTCACTCAAGTG	1200
Db	7656	TAGCAGAGATGGGGTTTTTACCATATGTTGCCAGAGGCTGGTCTCAAACTCTCTCAAGTG	7715
Qy	1201	ATTCGCCGCTCAGTCTCCCAAAAGTCTGGAAATTATAGGCGTGAAGTCACTGGGCTGGC	1260
Db	7716	ATTCGCCGCTCAGTCTCCCAAAAGTCTGGAAATTATAGGCGTGAAGTCACTGGGCTGGC	7775
Qy	1261	CGATTACTGTCTATTTTTCTTTATTTGCTATATATCCCAAGATCTAGACAGTGTGACATAT	1320
Db	7776	CGATTACTGTCTATTTTTCTTTATTTGCTATATATCCCAAGATCTAGAGCAGTGTGACATAT	7835
Qy	1321	AGTAGGTGCTCAATPAATPAATATGATGATGACAGGCTAGATATATACTTTCTTTTCTT	1380
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Qy	1381	TTTTTAAACATCTGACAACTTGGAGAAATAAATCAATCTGTGACTTGGCTTTTTC	1440
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Qy	1441	CTTATACCTTGTTATGACTTTTTCATATTTGCTCAACCTTATTTGTACTGTTTTTC	1500
Db	7956	CTTATACCTTGTTATGACTTTTTCATATTTGCTCAACCTTATTTGTACTGTTTTTC	8015
Qy	1501	ATTGTATCTATTTTATGACTGTAATTAATATGGCTTAAATTTGCTTATACATCTCTGCTC	1560
Db	8016	ATTGTATCTATTTTATGACTGTAATTAATATGGCTTAAATTTGCTTATACATCTCTGCTC	8075
Qy	1561	CACCTTAAAGGCCAAATTTTCAATCTGATGAAAGCTATGAACCCCTCCCGAGAGAA	1620
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Qy	1621	TACACACACACACACACTCACAACAAGTTTTTTTTTAATGTTTGGCACTAGACACAGA	1680
Db	8136	TACACACACACACACACTCACAACAAGTTTTTTTTTAATGTTTGGCACTAGACACAGA	8195
Qy	1681	AACCTGCATTAAGAGATGTTTTGTCATATTAATTAATAATACTAGTTGGGCAACGTGA	1740
Db	8196	AACCTGCATTAAGAGATGTTTTGTCATATTAATTAATAATACTAGTTGGGCAACGTGA	8255
Qy	1741	CTCAAGCCTGTACACACAGTACTTTGGAAAGTCCAAAGGTGGGTGATCATCTTGAGGTGAG	1800
Db	8256	CTCAAGCCTGTACACACAGTACTTTGGAAAGTCCAAAGGTGGGTGATCATCTTGAGGTGAG	8315
Qy	1801	AGTTGAGACCAAGCTGTGCAATATATGTAAGAACCTATCTCTATCAAAAATAACAAAATT	1860
Db	8316	AGTTGAGACCAAGCTGTGCAATATATGTAAGAACCTATCTCTATCAAAAATAACAAAATT	8375
Qy	1861	AGCTGGGTGTATGTATGTATGTATGATGCTGCTAGTCTCAAGTACTGGGAGGCTGAGGCAAGAA	1920
Db	8376	AGCTGGGTGTATGTATGTATGTATGATGCTGCTCAAGTACTGGGAGGCTGAGGCAAGAA	8435
Qy	1921	TTGCTTGAACCTGGAGGCAAGGTTTGACATGAGCCGAGATCCCAACCATCTGACATCCAGC	1980
Db	8436	TTGCTTGAACCTGGAGGCAAGGTTTGACATGAGCCGAGATCCCAACCATCTGACATCCAGC	8495
Qy	1981	CTGGGCGACACAGGAGACTCTATCTCAAAAATAATTAATTAATAATAAGATTCGGAG	2040
Db	8496	CTGGGCGACACAGGAGACTCTATCTCAAAAATAATTAATTAATAATAAGATTCGGAG	8555
Qy	2041	AGAAAACAAAACTAATTAAGATTTCTGTAGAGGTAGCAGAGATAGTAATTAATATATATATA	2100
Db	8556	AGAAAACAAAACTAATTAAGATTTCTGTAGAGGTAGCAGAGATAGTAATTAATATATATATA	8615
Qy	2101	AGTTTAATGCAATTTTAACTGTATCTTATTTGTTTATTTGGTTTATAAAGTAAACAAGC	2160
Db	8616	AGTTTAATGCAATTTTAACTGTATCTTATTTGTTTATTTGGTTTATAAAGTAAACAAGC	8675
Qy	2161	CAAAAAGTATGCAACTTCAAAACACCTACATPAATATCTATATATGAGAAAGTGAAGCACTT	2220
Db	8676	CAAAAAGTATGCAACTTCAAAACACCTACATPAATATCTATATATGAGAAAGTGAAGCACTT	8735

PA (SAGR-) SAGRES DISCOVERY.

XX Morris DW;

XX WPI: 2003-328604/31.

XX Recombinant nucleic acid useful for diagnosis and treatment of carcinoma
PT comprises a nucleotide sequence.

XX Claim 1; SEQ ID NO 100; opp; English.

XX The present invention relates to novel DNA and protein sequences which
CC are associated with carcinomas. The sequences are useful for: (i) for
CC screening drug candidates; (ii) for screening of bioactive agent capable
CC of binding to Carcinoma Associated Protein (CAP); (iii) for screening of
CC a bioactive agent capable of modulating the activity of CAP; (iv) for
CC evaluating the effect of a candidate carcinoma drug; (v) for diagnosing
CC carcinoma; (vi) for inhibiting the activity of CAP; (ix) as a biochip;
CC (x) for diagnosing carcinoma or a propensity to carcinoma; and (xi) for
CC determining Carcinoma Associated (CA) gene copy number. In addition, the
CC CA genes are useful as DNA vaccines and the CAP are useful as markers of
CC carcinoma including lymphoma. The present sequence is one such CA coding
CC sequence. Note: This patent is an equivalent to basic patent
CC US2002182586A1, for which no sequence data was published

XX Sequence 160482 BP; 44060 A; 32143 C; 33530 G; 49875 T; 0 U; 874 Other;

Query Match 17.4%; Score 434.2; DB 11; Length 160482;

Best Local Similarity 60.2%; Pred. No. 2.7e-79;

Matches 888; Conservative 1; Mismatches 525; Indels 61; Gaps 8;

Qy 606 TTTTGTGTTGTTAGAGACAGGCTGCTCTGACACCCAGGATGAGCAGATGG 665
Db 5181 TGTGTTTGTGTTCTGAGTAACGAGCTTGCTCTGTCGCCAGGCTGGAGTGAAGGG 5122
Qy 666 TGAACCATAGTGTGACGACGCTCAACCTCTGAGCTCAAGGATGCTGACCTGACG 725
Db 5121 CATATCTCGGCTCATGTAACCTCTGCTCTGCTGCTCAAGTATTCCTGCTCAGA 5062
Qy 726 CTCCCAAGTACCTGGAGCTACGAGCGTGCACACCGCTGG--CTAATTAATAAATT 783
Db 5061 CTCCCAAGTACCTGGAGCTACGAGCGTGCACACCGCTGG--CTAATTAATTTGTAAT 5002
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Db 5001 TTTTGTGAGACG--AGGTTTGTACATGTTGCTAGGCTAGCTCAACTCTGACTCAA 4943
Qy 844 GCAATCTCTTACCTTGGCATCCCAAGTGTGGGATTACAGGGGTGAGCACCATGTGC 903
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Qy 904 GGTACTTATTTCTTTACATTCATCTTTCCATTAAGATGTAATCAAGAAACAGGA 963
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Qy 1084 CTGAGTACGTGAATTACAGCGTGCACACACCATGCTTGGCTAAATTTTGTATTTTAA 1143
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Qy 1384 TTAACAATCTTGACAACTTTGAGATAATAATCAATCTGCAATTCCTTTTCACTT 1443
Db 4440 CAGATCTCAAAATTCACAAATCAAAATTCACATATATTTCCATAAATTCGCCCT 4381
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Db 4380 TTCTCTGTAATTTCTTACTGTTGCTTAATGATATCACTTAATTCAGGAAGAAATTCAAA 4321
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Db 4320 GTCACTTCAATTTTCTCTCTCTCAAGCAGATAAAGCACTAATTCAGAGCTCAAC 4261
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Db 4260 TTAACAGTCTCTCGAATGATGATGAGTTAACAGTTAACGACACAGACCAAACTAG 4201
Qy 1616 AGAATATCACACACACACACACTCAACACAGTTTCTTTTAAATTTTGAACCTAAGA 1675
Db 4200 GCACAGAGAGAGGGTGTTCATATTAATTCATCTCTGAGAGCCCTAAGGAATAGCT 4141
Qy 1676 CAAGAACTCGCATTAAGAGATGTTGTCAATTAATTAATAAT-----AATCAGTT 1729
Db 4140 CCAATTTTATAGAGAGAGAACTGAGGTTCCAGAAATTAAGAAATTCACACAGAGGCT 4081
Qy 1730 GGGCAGAGTACCTCAAGCTGTAAACACAGTACTTTTGAAGTCCAGAGTGGTGGATCAC 1789
Db 4080 GGGTGTGTGGCTCAGCTGATATCTGATACACTTTTGGAGGCCAAGCGGCGAGATCAC 4021
Qy 1790 TTGAGTGAAGATTTGAGACACAGCTGTGCTAATATGTGAAACCTATCTTACTAATA 1849
Db 4020 CTGAGGTCAAGAGTCCAGACAGCTGACAAACATGTGAAACCCCATCTTACTAATA 3961
Qy 1850 ATCAAAATTTAGTGTGTGTGATGATGATGCTGTGATCCAGCTACTCGGAGGCTG 1909
Db 3960 ATCAAAATTTAGTGTGTGTGATGATGATGCTGTGATCCAGCTACTTGTGAGGCTG 3901
Qy 1910 AGGCAAGAAATTTGTTGAACCTGAGAGGAGGAGTTCAGTGAAGC-----GAGATCC 1963
Db 3900 ACGAGAGAAATTTGTTGAACCTGAGAGGAGGAGTTCAGTGAAGGAGATGCG 3851
Qy 1964 CACCACTGCACTCAGCTGGGCGACACAGGAGACTCTATCTCAAAAAATTAATAAT 2023
Db 3840 TGCCACTGCACTCAGCTGGGCGACACAGGAGGAGACTCTCTCAAAAAATTAATAAT 3781
Qy 2024 AAAATTAAGGATGGAGAGAAACAAAATAATAG 2058
Db 3780 AGAAG 3746

RESULT 7

ACM4410/C
ID ACM4410 standard; DNA; 91760 BP.

AC ACM4410;

DT 18-NOV-2004 (first entry)

DE Human genomic sequence hnc41365.

XX Cytostatic; carcinoma; lymphoma; cancer; human; gene; ss.

OS Homo sapiens.

XX MO2003073826-A2.
XX 12-SEP-2003.
XX 28-FEB-2003; 2003WO-US006235.
XX 01-MAR-2002; 2002US-00087192.
XX (SAGR-) SAGRES DISCOVERY.
XX Morris DW;
XX WPI: 2003-328604/31.
XX Recombinant nucleic acid useful for diagnosis and treatment of carcinoma
XX comprises a nucleotide sequence.
XX
XX Claim 1; SEQ ID NO 844; Opp; English.
XX
XX The present invention relates to novel DNA and protein sequences which
XX are associated with carcinomas. The sequences are useful for: (i) for
XX screening drug candidates; (ii) for screening of bioactive agent capable
XX of binding to Carcinoma Associated Protein (CAP); (iii) for screening of
XX a bioactive agent capable of modulating the activity of CAP; (iv) for
XX evaluating the effect of a candidate carcinoma drug; (v) for diagnosing
XX carcinoma; (vi) for inhibiting the activity of CAP; (vii) for treating
XX carcinoma; (viii) for neutralizing the effect of CAP; (ix) as a biobchip;
XX (x) for diagnosing carcinoma or a propensity to carcinoma; and (xi) for
XX determining Carcinoma Associated (CA) gene copy number. In addition, the
XX CA genes are useful as DNA vaccines and the CAP are useful as markers of
XX carcinoma including lymphoma. The present sequence is one such CA coding
XX sequence. Note: This patent is an equivalent to basic patent
XX US2002182386A1, for which no sequence data was published
XX
SQ Sequence 91760 BP; 21177 A; 23589 C; 24129 G; 22845 T; 0 U; 20 Other;
Query Match 17.3%; Score 433.2; DB 11; Length 91760;
Best Local Similarity 61.6%; Pred. No. 3.8e-79;
Matches 906; Conservative 1; Mismatches 474; Indels 89; Gaps 10;
QY 606 TTTTGTGTTGTTTGAAGACAGGGTCTGCTGTGACCCAGGCGATGACAGAGTGG 665
DB TTTTGTGTTTAAATTAAGACGTGGTTTGTCTGTGCGCCAGGATGAGTACAGTGG 5507
QY 666 TGCACCATAGTCTACTGACGCTCACTCTGAGCTAAGGATCTGCTACCTCAGC 725
DB 5506 CATGATCATGCTCACTACACCTCTGCTCTGGGCTCAGGCAATCTCCACCTCAGC 5447
QY 726 CTCCCAAGTAGCTGGGACTACAGAGGTGACACCAACGCTGATTAATAAAT-TT 784
DB 5446 CT-CGAGTAGCTGGAACCTACAGGTGAGACACACACCTGCTTAATTTTATTTAT 5388
QY 785 TTTTGTAGAGCTGGGTCTTACTACGTGGCAGGGTCTTAACTCTGCTGCTCAAG 844
DB 5387 TGTGTGAGAGAGGGTCTCGTATGT-----GGCTGGTCTTGAACCTCTGGGCTCAAG 5334
QY 845 CAATCTCTTACTTGGCATCTCCAAAGTCTGGGATTTACAGGGGTGACCAACATGTGG 904
DB 5333 TGAATCTCCACCTCGGCTCTCAAGCGCTGGGATTTCAAGGTGTCAGGCTCTACACCCA 5274
QY 905 GGTACTTATTTCTTT-----ACATTCATCTTTCC 934
DB 5273 GCTCATTAATATCTTTTAAATCCAGATTAATCTGCTGCTGCCCCAAAGAACTTTT 5214
QY 935 AATAGATGTAAAGATCCACAGAACAGGATTAAGTCTTAATTTCTTCTTTTGTGA 994
DB 5213 ACTGTCTCCCAATCTTAAGATAGTGAATAAG---ACTGATTTTCTTTTGTGA 5157
QY 995 GACAGAGTCTCACTTATCACTCAACCTCCGTTG-----CTCACTGCA 1040
DB 5156 GGTGAGTAGTGTGTCTGTGCTTGAAGTGCAGGTGTGATCTTGAACCTCACTGCA 5097

QY 1041 ACTGTGCTCTCCCGGGTTCAAGYATTTCTCTGCTAAGCTCTCTGAGTACGTAATTA 1100
DB ACCCTCTCTCTTCTGCGTTTCAAGGATTTCTCTGCTCAGGCTCCAGAGTAGTGGACTG 5097
QY 1101 CAAGCTGACACCAACATGCTTGGCTAATTTTGTATTTTATAGAGAGATGGGGTTTAC 1160
DB 5036 CAGGTGGGCTCCACAGGCCCACTAAATTTTAAATTTTAAAGATAGGGGTTTCAC 4977
QY 1161 CATGTGCCAGGCTGGGTCTCAACTCTGACCTCAAGTATGCTGCTCAGTCTCC 1220
DB 4976 CACTTTGCGAGGCTGGTCTGAACTCTGACCTCAAGTATTCACCTGCTCCAGCTCC 4917
QY 1221 CAAAGTGTGAATTAATAGCGGTGATGATCTGCTGCGCGATTAATCTTATTTCTT 1280
DB 4916 CAAAGCGGTGGGATTAATAGGATGAGCCAAATGATCTGGCTGAAGATTAATTTCTTC 4857
QY 1281 TATTGCTAATCCCAAGATCTAGAGCACTGTCTGACATATATAGTGGTCTCAATTA 1340
DB 4856 TTTTGTAGCATTTTCTCTGTGATTAACGCA-----TGTGTGTACATTA 4810
QY 1341 TTGATGAATGACAGCGCTAGATATAACTTTCTTTTCTTTTAAACAATCTTGACA 1400
DB 4809 TGTTTTCTGCTGATTTTGAACCTGTGACTTTCTTTCTTCTTCTTCTTCTTCT 4750
QY 1401 ACTTTCAGAAATTAATACATCTTGATTCGCTTTTCTACTTATACACTTGTATGACT 1460
DB 4749 CTCTCTCTTCTT--CTCTCTCTTCTTCTCTCTCTCTCTCTCTCTCTCTCTCT 4692
QY 1461 TTTTCAATTTGCTTCAACCTTTATGTTAGTCTTTTTCATGTTTATTTATTTAGTAC 1520
DB 4691 TTTCT 4632
QY 1521 TGAATTAATAGCTTAATTTGCTTAATACATCTCTGCTCCACTTATAGAGCCAAATTT 1580
DB 4631 CTCTCTCTTCT 4584
QY 1581 ACAAATCTGATGAAGAACTATGAAACCTCTCCCAAGAAATACACACACACACACT 1640
DB 4583 CTTCT 4524
QY 1641 CACACACAGTTTTTTTAAATGTTTGAACCTTAAGCAAGAAACCTGCACTTGAAGAT 1700
DB 4523 TCTTCTCTTCT 4471
QY 1701 TGTTCATTAATTAATTAATTAATTAATTAATTAATTAATTAATTAATTAATTAAT 1760
DB 4470 CTTTTCTTTTAAATTAATTAATTAATTAATTAATTAATTAATTAATTAATTAAT 4411
QY 1761 ACTTGAAGTCCAAAGTGGGTGATCACTTGAAGTGAAGATTCGAACCAAGCTGTGTC 1820
DB 4410 ACTTTGGAGGCGCAAGGGGGGGGATCACTTGAAGTCAAGTTCACAGCTCAGCTGCGC 4351
QY 1821 AATATGAGAAACCTTAATCTTAATTAATTAATTAATTAATTAATTAATTAATTAAT 1880
DB 4350 AACATGTGAAACCTGAGTCTTAATTAATTAATTAATTAATTAATTAATTAATTAAT 4291
QY 1881 GCTGTAGTCCAGCTAATCTGGAAGTGAAGGCAAGAAATTTGTTGAACCTGGAAGGCA 1940
DB 4290 GCTCTTAATCCAGCTAATCTTGAAGGCTGAGGCAAGAAATTCCTTGAACCAAGAGGG 4231
QY 1941 GAGGTGAGTAGGCGGAGATCCACCACTGACCTCCAGCTGGGCGACACAGGAGACT 2000
DB 4230 GAGGTGAGTAGGCGGAGATCCACCACTGACCTCCAGCTGGGCGATATATAGAGAAAT 4171
QY 2001 CTATCTCAAAAAATTAATTAATTAATTAATTAATTAATTAATTAATTAATTAAT 2030
DB 4170 CCATCTCAAAATTAATTAATTAATTAATTAATTAATTAATTAATTAATTAATTA 4141
RESULT 8
ABD3357/c
ID ABD3357 standard; DNA; 38538 BP.
XX

AC ABD33357;
XX 18-NOV-2004 (first entry)
XX
DE Human cancer-associated (CA) gene HD07-065.
XX
XX Human cancer-associated protein; CAP; cancer-associated gene; CA; gene;
XX
XX Human cancer-associated protein; CAP; cancer-associated gene; CA; gene;
XX
XX ds; cancer; cytostatic.
XX
XX Homo sapiens.
XX
XX WO2004058146-A2.
XX
XX 15-JUL-2004.
XX
XX 15-DEC-2003; 2003WO-US040081.
XX
XX 17-DEC-2002; 2002US-00322281.
XX
XX (SAGR-) SAGRES DISCOVERY INC.
XX
XX Morris DW, Malandro MS;
XX
XX WPI; 2004-499109/47.
XX
XX Novel human cancer associated protein encoded within open reading frame
XX of cancer associated gene, useful as targets for diagnosing cancer.
XX
XX Claim 16; SEQ ID NO 448; 182bp; English.
XX
XX The invention relates to cancer-associated proteins (CAP) and the cancer-
XX associated (CA) nucleic acids encoding them. The invention also relates
XX to a method for treating cancers involving administering to a patient an
XX inhibitor of CAP, and a method of screening for anticancer activity in a
XX potential drug involving providing a cell that expresses a CA gene,
XX contacting a tissue sample derived from a cancer cell with an anticancer
XX drug candidate and monitoring the effect of the anticancer drug candidate
XX on expression of the CA gene. The CAP proteins are useful for detecting
XX cancer associated with expression of a CAP protein in a test cell sample
XX and for screening for a bioactive agent capable of modulating the
XX activity of a CAP protein. The CA nucleic acids are useful for diagnosing
XX cancer, involving determining the expression of a CA nucleic acid in a
XX tissue. This sequence represents a human CA gene of the invention. Note:
XX The sequence data for this patent did not form part of the printed
XX specification, but was obtained in electronic format directly from WIPO
XX at ftp.wipo.int/pub/published_pct_sequences
XX
XX Sequence 38538 BP; 9418 A; 9893 C; 9523 G; 9684 T; 0 U; 20 Other;
SQ
Query Match 17.0%; Score 425.4; DB 13; Length 38538;
Best Local Similarity 60.0%; Pred. No. 1.3e-77;
Matches 879; Conservative 1; Mismatches 552; Indels 33; Gaps 9;
QY 567 CCAAGATCACACAGCTTGAAGTGTGACAGTTGGGTTTTTTTGTGTTTGAAG 626
DB 35304 CAAAAAATTAATAATATTTTGTGTTTGAATATTAAGGCGCTTTTGGAG 35245
QY 627 ACAGGGCTTCTC-TGTCAACCCAGGATGACAGTGTGCAACCATAGGTCACTGCA 685
DB 35244 ATGAAGTTTGTCTTGTGTCAGGCTGAGTGCATGTGTATCTTGTGCTCACAGCA 35185
QY 686 GCTCAACCTCTAGCTCAAGGATCTGACCTGACCTCCCAAGTAGCTGGACTTA 745
DB 35184 ACCTTCACCTTCAGAGGTTCAAGCAATTCCTCTCTCAAGCTCTGAGTACCTGGATTA 35125
QY 746 CGAGCGTGACACACACGCTGAGCTTAATTAATAATTTTGTGAGAGCTGGGTCTTA 805
DB 35124 CAGGCAATGCGGCACACACCGCTAATTTTGA---TTTATGAGAGAGGGGTTTCT 35068
QY 806 CTAGCTGGCCAGGCTGTCTTAATCTCTGGCTTCAAGCAATCTCTCACTTGGCATC 865
DB 35067 CCAATGTTGTGTCAGGCTGGTCTTGAATCTTCGACCTCAGGTATCAACCTCTTGGCTTC 35008

QY 866 CCAAGTGTGGATTAACAGGGTGAAGCCACATGTGCGCTACTTAATTTCTTACATTC 925
DB 35007 CCGAAGTTTGGATTAACAGGCAATGACCACTGGCGCTGGC-----CTAGTTAAGGA 34954
QY 926 CATCTTTCCAATGATGATGATCAAGAACAGGATTAATCTGCTTAATTTCTTCTTT 985
DB 34953 CTTTTCATATTAATTAATTAATTAATTAATTAATTAATTAATTAATTAATTAATTA 34894
QY 986 CTTTTCGACAGAGCTGACCTTCACTCACTCAACCTCGGTCAGCTCACTGCACTC 1045
DB 34893 GTTTCGAGGCTGAGTGCATGACAGACC-----TTGGCTCACTGCACTC 34845
QY 1046 TGCTCCCGGGTTCAGAGTATCTCTGCTGCTTAAGCTCTGAGTAGTGAATTAACAGC 1105
DB 34844 CGCTCCCGGGTTCAGAGTATCTCTGCTGCTTCCGCTGCTCCAGAGTGGATTAACAGC 34785
QY 1106 GTGCAACAATGCTTGGCTTAATTTTGTATTTTGAAGAGAGAGAGGAGGTTTACATGT 1165
DB 34784 GCTGCAATCAATGCTTGGCTTAATTTTGTATTTTGAAGAGAGAGGAGGTTTACATTA 34726
QY 1166 TGCCAGGCTGCTGCTCAAACTCCGACCTCAAGATCTGCTGCTCAAGTCCCAAG 1225
DB 34725 TGCCAGGCTGCTGCTCAAACTCCGACCTCAAGATCTGCTGCTCAAGTCCCAAG 34666
QY 1226 TGCTGAATTAATGAGCTGAGTCACTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 1285
DB 34665 TGCTGAGATGAGAGTGTGAGCCACACACACTCGGCTTTATTTATTTTTCAGACAGG 34606
QY 1286 CTATATCCCAAGATCTGAGCAGTGTGACATATAGTAGTGTCTCAATTAATTAATGAT 1345
DB 34605 GTATCTC-----TCTGTCGCCAGGCTGAGAGTCAAGTGTGAGTCAAGCTGCTGCA 34552
QY 1346 GAATGACAGCTGATATTAATCTTTCTTTCTTTTAAACATCTTGACACTTT 1405
DB 34551 GCTCAACCTCTGAGGCTGCAAGCA--ATCTCTGCTGCTGCTGCTGCTGCTGCTGCA 34494
QY 1406 GCAAAATTAATTAATTAATTAATTAATTAATTAATTAATTAATTAATTAATTAAT 1465
DB 34493 TACAGTGTCAATTAATTAATTAATTAATTAATTAATTAATTAATTAATTAATTAAT 34434
QY 1466 ATATGCTTCAAACTTTATTTGTTTACGTTTTCATGTTTACTATTTTACTGAT 1525
DB 34433 CTATATGCTTCAAACTTTATTTGTTTACGTTTTCATGTTTACTATTTTACTGAT 34374
QY 1526 AATATGCTTCAATTTGCTTATTAATCTCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 1585
DB 34373 AATATGCTTCAATTTGCTTATTAATCTCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 34314
QY 1586 TCGATGAAGCTATGAACCTCTCCCGAGAGAAATACACACACACACACTCAAC 1645
DB 34313 CTTTCTGCTTCT 34254
QY 1646 ACAATTTTTTTTAAATGTTGCACTTAAGCAAGAAACCTGCAATGAGATGTTTGTTC 1705
DB 34253 CTGATATGCTGATTAATCTGCAATATGATATTAATTAATTAATTAATTAATTAATTA 34194
QY 1706 ATA--TTAATTAATAATTAATTAATTAATTAATTAATTAATTAATTAATTAATTA 1763
DB 34193 TTCTTTGTTTAAATTAATTAATTAATTAATTAATTAATTAATTAATTAATTAATTA 34134
QY 1764 TTGGAATGCAAGGTTGGTGGATCACTTGAAGTGAAGATTTGAGACCAAGCTGCTCAAT 1823
DB 34133 TTGGAGGCTGTGTGTGGGCGGATCACTTGAAGTGAAGATTTGAGACCAAGCTGCTCAAT 34074
QY 1824 ATGATGAACCTTAATCTTAATAATAATAATAATAATAATAATAATAATAATAATAATA 1883
DB 34073 ATGATGAACCTTAATCTTAATAATAATAATAATAATAATAATAATAATAATAATAATA 34014
QY 1884 TGTATGCTGAGCTATCTGAGAGCTGAGGCAAGAAATTTCTTGAATCTGAGAGGCGAG 1943
DB 34013 TGTATGCTGAGCTATCTGAGAGCTGAGGCAAGAAATTTCTTGAATCTGAGAGGCGAG 33954
QY 1944 GTTGCAGTGAAGCGAGATCCACCACTGACCTCGAGGCGACACAGAGACTCTA 2003

D6	33953	GTTGCACTGGAGCCAAATGACACACTATTGCCTCAGCACAGGCGACA-AGAAGACTCTG	338955
Oy	2004	TCTCAAAAAATAAATAATAATAAT	2028
D6	33894	TCTCAAAAAAAAAAAAAAAAAAGCAT	33870
 RESULT 9			
ID	ABK83569	standard; cDNA; 122888 BP.	
AC	ABK83569;		
XX			
DT	14-AUG-2002	(first entry)	
XX			
DE	Human CDNA differentially expressed in granulocytic cells #140.		
XX			
KM	Human; ss; granulocytic cell; DNA chip; bacterial infection;		
KM	viral infection; parasitic infection; protozoal infection;		
KM	fungal infection; sterile inflammatory disease; psoriasis;		
KM	rheumatoid arthritis; glomerulonephritis; asthma; thrombosis;		
KM	cardiac reperfusion injury; renal reperfusion injury; ARDS;		
KM	adult respiratory distress syndrome; inflammatory bowel disease;		
KM	Crohn's disease; ulcerative colitis; periodontal disease;		
KM	granulocyte activation; chronic inflammation; allergy.		
XX			
OS	Homo sapiens.		
XX			
PN	WO200228999-A2.		
XX			
PD	11-APR-2002.		
XX			
PF	03-OCT-2001; 2001WO-US030821.		
XX			
PR	03-OCT-2000; 2000US-0237189P.		
XX			
PA	(GENE-) GENE LOGIC INC.		
PI	Beazer-Barclay Y., Weisman SM, Yamaga S, Vockley J;		
PT	WPI, 2002-435328/46.		
DR			
XX			
PS	Claim 1; SEQ ID NO 140; 114pp; English.		
XX			
CC	The invention relates to detecting (M1) granulocyte (GC) activation		
CC	(GCA), by detecting the level of expression of gene(s) (Gs) identified by		
CC	DNA chip analysis as given in the specification, and comparing the		
CC	expression level to an expression level in an unactivated GC, where		
CC	differential expression of Gs is indicative of GCA. Also included are		
CC	modulating (M2) GA by contacting GC with an agent that alters the		
CC	expression of at least one gene in Gs; (2) screening (M3) for an agent		
CC	capable of modulating GCA or an inflammation (especially chronic) in a		
CC	tissue, an allergic response in a subject, exposure of a subject to a		
CC	pathogen or sterile inflammatory disease using the gene expression		
CC	profile; (3) detecting (M4) an inflammation (especially chronic) in a		
CC	tissue, an allergic response in a subject, exposure of a subject to a		
CC	pathogen or sterile inflammatory disease, by detecting the level of		
CC	expression in a sample of the tissue of gene(s) from Gs, where the level		
CC	of expression of the gene is indicative of inflammation; (4) treating		
CC	(M5) an inflammation (especially chronic) or in a tissue, an allergic		
CC	response in a subject, exposure of a subject to a pathogen or sterile		
CC	inflammatory disease, by contacting a tissue having inflammation with an		
CC	agent that modulates the expression of gene(s) from Gs in the tissue. M1		
CC	is useful for detecting GCA; M2 is useful for modulating GA; M3 is useful		
CC	for screening an agent capable of modulating GCA preferably in an		
CC	inflammation in a tissue; M4 is useful for detecting an inflammation		
CC	(especially chronic) in a tissue, an allergic response in a subject,		
CC	exposure of a subject to a pathogen or sterile inflammatory disease (e.g.		

	CC	pneumonia, rheumatoid arthritis, glomerulonephritis, asplenia, thrombocytopenia,
	CC	cardiac reperfusion injury, renal reperfusion injury, ARDS, adult
	CC	respiratory distress syndrome, inflammatory bowel disease, Crohn's
	CC	disease, ulcerative colitis, periodontal disease; also bacterial
	CC	infection, viral infection, parasitic infection, protozoal infection,
	CC	fungal infection and MS is useful for treating one of the above
	CC	conditions. The present sequence represents a gene differentially
	CC	expressed in granulocytes. Note: The sequence data for this patent did
	CC	not form part of the printed specification, but was obtained in
	CC	electronic format directly from WIPO at
	XX	ftp.wipo.int/pub/published_jct_sequences
SQ	Sequence 122888 BP; 28761 A; 33410 C; 31919 G; 28798 T; 0 U; 0 Other;	
	Query Match	16.7%; Score 417.2; DB 6; Length 122888;
	Best Local Similarity	61.1%; Pred. No. 8.1e-76;
	Matches 894; Conservative	0; Mismatches 503; Indels 65; Gaps 11;
QY	604	TTTTTTTGTGTTTGTTTAGAGACAGGGCTTCCTGCTGCACCAGCATATGACACAGT 663
DB	70248	TTTTTTTTTAAATTTTAAATTTTAAACACAGAGTCCTGCTGTGTCACCAAGCTGAGTGACGT 70307
QY	664	GATGCAACCATATGATCAGCTGACAGCCCTCAACCTCTGAGCTCAAGGAATCTGTCATCA 723
DB	70308	GTGTCAAACAACACTGACGACGACACTCAAAC-CCTGGGCTCAAGCAATCTTCCACCTCA 70366
QY	724	GCTCCCAAGTATCTGGAGATCAGACCGTGACCAACCAACGCTGCTA----- 772
DB	70367	GCTTCCCAGAATCTGGGACATAAGGACATATGACACCAATGCCACATATTATTTTTTTT 70426
QY	773	-----TTAAAAATTTTTTTTGTAGAGACTGGGCTCTACTAGTGGCACAGGCTGTCT 827
DB	70427	TTTACTTTTAAAATTTTTTTGTAGAAATGGAGCTCCTACTTAATATGCTTAGCTGCTCTC 70486
QY	828	AAATCTCTGCTTCAAGCAATCTCTCTACCTTGGCATCCCAAGCTGAGATTACAGGG 887
DB	70487	AAATTTCTTGGGCTTAAGCAATCTCTGCTGGGCTCCCAAGTTGGGATTAAGT 70546
QY	888	GTGAGCAACCATGTGGGCTAATTATTTCTTATCATCTCTTCCATAGATATAG 947
DB	70547	GTG----ACCATGTCAGCCAGCCTTAATTTCTAATATGTAAGTATTAATAGCATTTGG 70602
QY	948	ATCCACGAACAGGAGATTATGCTCTATTTTCTCTCTTTTGTAGACAGACTGCAC 1007
DB	70603	GGATGCTCAATAT--TTGCTTCTCTTTTTTTTTTTTTTGTAGATGGAGCTTAGC 70657
QY	1008	TTCAATCACTCAACCTCCG-----TTGAGCTCACTGCAACCTCTGCTCC 1053
DB	70658	ACTGTCACTTAGCTGAGTGCATATGGCGGATTTCAATTTCACTGCAACTCTGCTCCT 70717
QY	1054	GAGTTCAAGATTTCTCTGCTCTGAGCTCTCTGAGTAGTGCATTAACAAGCTGCACCA 1113
DB	70718	GGGTTTCAAGAGATTCTGTGCTCTCAAGCTCTCTAGTAGTGCAGGACTACAGCACCGCA 70777
QY	1114	CCATGCTTGGCTAATTTTTTGTATTTTATAGACAGATGGGGTTTACATGTTGCCACAG 1173
DB	70778	CCAATCCGGCTAATTTTTTGTATTTTATAGATAGATAGAGTGTGCTATGTTGGCCAGG 70837
QY	1174	CTGGTCTCAAACTCTGACCTCAAGATATGCTGCTCTCACTCTCCAACTGCTGGA 1233
DB	70838	CTGGTCTCAAACTCTTCACTTC--GTGATCCACATCTCTCAAGCTCCCAAACTGCTGGA 70895
QY	1234	TTATAGCGGTAGTCACTGTGCTGGCCGATTAATCTGTATTTTCTTATTTGCTATATCC 1293
DB	70896	TTACAGGTGTAGCACCAACATCTCG-----CATATATTTTAAAGTATTAATA 70944
QY	1294	COAGATCTAGACAGTGTCTGACATATATAGTGTCTCAATTAATATGATGATGCAC 1353
DB	70945	ATGTCCAGAGCCAAAAGTCAAGAACCACTCTTAAATACATAAATATGAT-----ATC 70999
QY	1354	AGCGTAGATTAATCTTTCTTTCTTTTAAACAAATCTTGAACAATTTGACAGATA 1413
DB	71000	AATCATTTATTTATTTTCAAAATTAATCTTTTAAATAATTTCAACAGAGCATMAG-CTC 71058

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QY 1414 AATACATCTTGATCTGCTTTTCACTATACCTTGTATGACTTTTTCATATGSC 1473
DB 71059 CATGAGAGAGGTTTCTTTTCTTTTATTTATGAAAGTAACTCCCTAGAACAA 71118
QY 1474 TCAACCTTATTTGTTACTGTTTTTCACTTTTACTATTTTATGCTAGTAATATATGSC 1533
DB 71119 TGCCGTGAACATATGAGGTGTTCAATTAATAGTATGAAATGAATTAATATTTT 71178
QY 1534 TTAATTGCTTATATCATCTCTCTGCTCCACTTTAAGGCCAAATTTTACAAATCTGATGA 1593
DB 71179 GGGTTTACCCATTTCTGATACCTGAGCCTCAAACTATCTGCTGTTCAATCTATTA 71238
QY 1594 AAGTATATACCCCTCTCCAGAGAAATACACACACACACACACACACAGTTT 1653
DB 71239 TTCCTGAAGATGAATAATATACATTAATATTTATTAAGAACATTTGTTCTATATTTT 71298
QY 1654 TTTTAA---TGTTGCACTAAGACAAAGAACTGATTAAGAGATGTTGTTCAATAT 1709
DB 71299 TCATGACATTTTATATACAGAGTAAGATTTCTCAGGGTATATATACCTAAGAAATGAAC 71358
QY 1710 TAATTAATAAATTAATCACTGTTGGGCACTGATGCTCAAGCTGTAACTCAAGTATTTGAA 1769
DB 71359 TGTAGTAAATTAATCTGCGCAGGCAAGGTGCTCATGCTGTATATCCAGACATTTGGGA 71418
QY 1770 GTCCAAAGGTGGGTGATCACTTGAAGTGAAGATTCAGAGACAGCCTGCTCAATATGGTG 1829
DB 71419 GCGCGAGGCAAGTGAATCAGC--AGGTACAGGGGATCAAGACCATCTGCGCCACACATG 71476
QY 1830 AAACCTATCTCTATAAATAATCAAAATAATAGTGGGTATAGTATGATGCTGTAAT 1889
DB 71477 AAACCTCTCTCTATAAATAATCAAAATAATAGCAGAGTGGGGGCGTGCGCTGTAGT 71536
QY 1890 CCCGCTACTGCGGAGGCTGAGGCAAGAAATGCTTGAACCTGGAGGCAAGGTTGCA 1949
DB 71537 CCGAGCTAAGTGAAGGCTGAGGCAAGAAATGCTTGAACCCAGAGAGTGAAGGTTGCA 71596
QY 1950 GTGAGCCGAGATCCACACCTGCACTCCAGCCTGGGCGACACAGGAGACTCTATCTCAA 2009
DB 71597 GTGAGCCGAGATCCACACCTGCACTCCAGCCTGGGCTGACAGAGCAAGATCTCATCTCAC 71656
QY 2010 AAAAAATAAATAAATAAATAA 2031
DB 71657 AAACCAAAAAGAAAAAAA 71678

RESULT 10
AAF97850/C
ID AAF97850 standard; DNA; 6405 BP.
XX
AC AAF97850;
XX
DT 31-MAY-2001 (first entry)
XX
DE Human neuroblastoma cell line NB-1 Ip36 nucleotide sequence SEQ ID NO:64.
XX
KM Human; chromosome 1; Ip36; neuroblastoma cell line; NB-1; anticancer;
KM tumour suppressor; human Ip36 homozygosity deletion domain; tumour;
KM diagnosis; db.
XX
OS Homo sapiens.
XX
PN WO200116311-A1.
XX
PD 08-MAR-2001.
XX
PF 31-AUG-2000; 2000WO-JP005930.
XX
PR 31-AUG-1999; 99JP-00245962.
PR 09-MAY-2000; 2000JP-00136266.
XX
PA (HISM) HISAMITSU PHARM CO LTD.
PA (CHIB-) CHIBA PREFECTURE.
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XX
PI Nakagawara A;
XX
DR WPI; 2001-226686/23.
XX
PT Human Ip36 homozygosity deletion domain from the 36-position of first
PT chromosome short arm in human neuroblastoma cell lines, applicable e.g.
PT in gene diagnosis of tumors as well as in developing anti-cancer drugs.
XX
PS Example 8; Page 93-95; 226pp; Japanese.
XX
CC The present invention describes a homozygosity deletion domain co-
CC existing in the 36-position of the first chromosome short arm (Ip36) in
CC human neuroblastoma. Also described are base sequences from the Ip36
CC position of human neuroblastoma cell lines (NB-1 and MAB-ND-SCH-1),
CC which are tumour suppressor genes in human neuroblastoma. The genes are
CC tumour suppressor genes, base sequence data of which are applicable as
CC tumour markers and reagents in studying mechanism of tumour body
CC formation, and gene diagnosis of tumours as well as in developing anti-
CC cancer drugs. AAF97878 to AAF97829 represent PCR primers used in the
CC exemplification of the present invention, and AAF97830 to AAF97874
CC represent sequences given in the exemplification of the present invention
XX
SQ Sequence 6405 BP; 1650 A; 1449 C; 1383 G; 1923 T; 0 U; 0 Other;

Query Match 16.7%; Score 416.2; DB 5; Length 6405;
Best Local Similarity 59.9%; Pred. No. 7e-76;
Matches 856; Conservative 1; Mismatches 504; Indels 68; Gaps 7;

QY 603 GTTTTGTGTTGTTGTTTGAAGACAGAGGCTTCTCTCTGTCACCCAGGACATGACAG 662
DB 3636 GTCTTATTTTATTTTATTTTGAAGTGAAGTTTCACTGTTGTCAGGCTGAGTGAA 3577
QY 663 TGTGCAACCATAGCTCACTGCAAGCTTCACTCTGAGCTCAAGAGATCTGCTGACTC 722
DB 3576 TGGCAGCATCTCAGCTTACAAACAACCTGCTCTCTGCTCAAGTCAAGTATCTCTGCTC 3517
QY 723 AGCTCCCAAGTATCTGAGGATACAGAGGTCACACACAGCCCTGGCTTAATTAATAAAT 782
DB 3516 AGCTCCCAAGTATCTGAGGATACAGAGGTCACACACAGCTCAAGTATTTTGTGA 3457
QY 783 TTTTGTGAGAGCTGGGCTCTTACTGATGAGCCAGGCTGCTTAAACTCTGAGCTTCA 842
DB 3456 TTTTGTGAGAG--TGGGTTTCTCCATGTTGTGTAAGGCTAGTCTCAAACTCCGACTCA 3398
QY 843 AGCAATCTCTTACCTTGGCATCCCAAGTGTGAGATTAAGGGGTGAGCCACATGTG 902
DB 3397 GGTGATCCGCTGCTGCGCTCCGCAAGTGTGAGATTAAGGGGTGAGCCACATGTG 3338
QY 903 CGGCTACTTATTTCTTTCATCTCATCTTTCATTAAGATTAAGATCCACAGAAAGG 962
DB 3337 TGGCTATTTTCTTACT----- 3322
QY 963 ATTACTGCTATTTCTTCTCTTTCTTTTGAAGACAGTCTCACTTCATCACTCAACC 1022
DB 3321 -TTTAAGATGAGTTTGTCTCTGTGTGTCAGAGTGAAGTGAATGAGCATGATC----- 3269
QY 1023 TCCGTTAGCTCACTGCAACCTTGTGCTCCCGGTTCAAGATTTCTGCTGAAGCT 1082
DB 3268 ----TCGGCTCACTGCAACCTCGCTCCAGGTTCAAGTATCTTGTGCTCAAGTCT 3214
QY 1083 CTTGAGTATGAGTGAATTAACAAGGTGACACACATGTTGGCTTAATTTTGTATTTTGA 1142
DB 3213 CCAAGTATGAGTGAATTAACAAGGTGACACACATGTTGGCTTAATTTTGTATTTTGA 3156
QY 1143 GCAAGATGAGGTTTATTAACAATGTTGCCAGAGTGTGTTCAAACTCTGACCTCAAGTAT 1202
DB 3155 GTAGAGATGAGGTTTCTCCATGTTGTTAGGCTGTGTTCAAACTCCCAACTC-AGTAT 3097
QY 1203 CTGCTGCTCAAGTCTCCCAAGTGTGAGATTAAGGCTGAGTCACTGTGCTGCGCG 1262
DB 3096 GCAACAGCTCTGCGCTCCCAAGTGTGAGATTAAGGCTGAGTCAAGGCTGAGGCTGCGCGCG 3037
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OY 1263 ATTACGTCTATTTCTTTATTTGCTATATCCAGATCTAGAGCAGTGTGACATATAG 1322
DB 3036 TAGGTATCTTTATATACAGATGTTATACAAATTTCTTTAGAACATCTTTAGAAATTA 2977
OY 1323 TAGGTGCTCATATAATATGATGAAATGACAGGCTGATATATACTTTCTTTCTTTT 1382
DB 2976 ATACTAGAAAAAAGGTTT-----CTCTGACAAATATATAGGCTTTCTTTTACGGA 2924
OY 1383 TTTAAACATCTTGACACACTTTGACAAATATATATCAATCTTGACTTGTCTTTTCACT 1442
DB 2923 GTGAAATTTGACAAATTTATCTTCCCTTTTGCTTTAAATTCATGTTCCAAATTAAT 2864
OY 1443 TATCACCTGTTATGACTTTTTCATATTTGCTCCAAACCTTATGTTACTGTTTTCAT 1502
DB 2863 TACAAATTAACCAATTTAGCCGAGGCACTATACATTTAGTTTCTTTCTTACATTAATTT 2804
OY 1503 TGTATCTATTTTATGCTGATGATATATATGCTTATATTTGCTTATACATCTCTGCTCCA 1562
DB 2803 TTACTTATTTTATTTTAAATAGCTTCACTGATATATTTTGTGTAAAGCTGCTCAATCTT 2744
OY 1563 CTTTGAAGGCAAAATTTACAAATCTGATGAAAGCTATGAAACCTCTCCCAAGAAATA 1622
DB 2743 TTTCTAGAAAGTAACTGATTAATTAATTAATTAATTAATTAATTAATTAATTAAT 2684
OY 1623 CACACACACACACACACCTC-ACACACAGTTTTTTTTTATGTTTGCATTAAGACAAATA 1681
DB 2683 TCTGCTATTTTAACTGTTAAATATGTTGTGTATGTTAAAGTTTCTTAAATTAATTAAT 2624
OY 1682 ACTGTCATTTAGAGATGTTTGTTCATATTTTAAATTAATTAATTAATTAATTAATTAAT 1741
DB 2623 AAAACCAACACACACACAGCAAGCTTTTAAAGTTTAAATTAATTAATTAATTAATTAAT 2564
OY 1742 TCAGCCCTGTAACCACTAGTCTTTGAAAGTCCAGAGTGTGTGATCACTTGAAGTGAATA 1801
DB 2563 TCACCCGTAATATCCCAACACTTTGTGGAGGCGCAAGCGGTGATCACTTGAAGTGAATA 2504
OY 1802 GTTGAGACCGAGCTGTGATATGTTGTAATGTTGTAATGTTGTAATGTTGTAATGTTGTAAT 1861
DB 2503 GTTCAAGACCGAGCTGTGATATGTTGTAATGTTGTAATGTTGTAATGTTGTAATGTTGTAAT 2444
OY 1862 GCTGAGTGTAGTATGATGATGCTGCTGTAATGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 1921
DB 2443 GCTGAGTGTAGTATGATGATGCTGCTGTAATGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 2384
OY 1922 TGTCTGAACCTGAGGAGGAGAGGTTGCTGATGAGCCGGAATCCCACTGCTGCTGCTGCTGCT 1981
DB 2383 CACTTGAACCCGAGGAGTGAAGTTGCTGATGAGCCAAAGGTCAACCACTGCTGCTGCTGCTGCT 2324
OY 1982 TGGGAGACACAGAGACTCTATCTCAAAAAATTAATTAATTAATTAATTAATTAATTAAT 2030
DB 2323 TGGATGACAGAGAGACTCTGCTCTCAAAAAATTAATTAATTAATTAATTAATTAATTAAT 2275

RESULT 11
ADP13332/c
ID ADP13332 strand: DNA; 126990 BP.
AC ADP13332;
DT 26-AUG-2004 (first entry)
DE Renal cell carcinoma differentially expressed gene #68.
XX de; diagnosis; non-blood disease; solid tumor; gene expression;
XX peripheral blood mononuclear cell; renal cell carcinoma; prostate cancer;
XX head/neck cancer; differential expression.
OS Homo sapiens.
XX
XX WO2004048933-A2.
XX
XX 10-JUN-2004.
XX
```

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PF 21-NOV-2003; 2003MO-US037481.
XX
PR 21-NOV-2002; 2002US-0427982P.
PR 03-APR-2003; 2003US-0459782P.
XX
XX (AMHP) WYETH.
XX (TWIN) TWINE N C.
XX (BURC) BURCZYNSKI M E.
XX (TREP) TREPICCHIO W L.
XX (DORN) DORNER A.
XX (STOV) STOVER J A.
XX (SLON) SLONI D K.
XX
XX Twine NC, Burczynski ME, Trepicchio WL, Dorner A, Stover JA;
XX Sloni DK;
XX
XX WPI; 2004-460799/43.
XX
XX Diagnosing non-blood disease such as solid tumor, involves comparing
XX differential expression profile of specific genes in peripheral blood
XX sample of subject with reference expression profile of specific genes.
XX
XX Disclosure; SEQ ID NO 68; 350pp; English.
XX
XX The invention relate to a method of diagnosing (M1) non-blood disease
XX such as solid tumor by providing peripheral blood sample of human having
XX non-blood disease, and comparing an expression profile of specific genes
XX in the peripheral blood sample to reference expression profile of the
XX genes, where each of the genes is differentially expressed in peripheral
XX blood mononuclear cells (PBMCs) of patients having the disease as
XX compared to PBMCs of normal humans. The method is useful for diagnosing
XX non-blood disease such as solid tumor. The solid tumor is chosen from
XX renal cell carcinoma (RCC), prostate cancer and head/neck cancer. The
XX peripheral blood sample comprises enriched PBMCs. The peripheral blood
XX sample is a whole blood sample (claimed). (M1) is useful for identifying
XX genes that are differentially expressed in peripheral blood samples
XX isolated at different stages of progression, development or treatment of
XX RCC and/or other solid tumors. This sequence corresponds to a gene that
XX is differentially expressed and detected by the method of the invention.
XX (Note: this sequence is not given as part of the printed specification
XX but was obtained from WPI in electronic format at
XX ftp://pub/published_pat_sequences).
XX
XX Sequence 126990 BP; 36683 A; 27389 C; 27065 G; 35853 T; 0 U; 0 Other;
XX
XX Query Match 16.6%; Score 416; DB 12; Length 126990;
XX Best Local Similarity 60.2%; Pred. No. 1.4e-75;
XX Matches 859; Conservative 1; Mismatches 536; Indels 32; Gaps 9;
XX
XX 611 TTGTTGTTGTTTAAAGACAGGAGTCTTGTCTGTGACCCGAGCATGACAGAGTGTGCAA 670
DB 27332 TTTCTTAATGCTGATGAGAGTCTCACTGTGACCCAGGCTGAGTGCATGTGTGGA 27273
OY 671 CCATAGTCACTGACGCTCAACCTCTGAGCTCAAGGAGTCTGTACCTTCAAGCTCCC 730
DB 27332 TTTCTTAATGCTGATGAGAGTCTCACTGTGACCCAGGCTGAGTGCATGTGTGGA 27273
OY 27272 TCTCA-----CCTCAACCTCCAGGTTCAAGGATTTCTTCTCCAGCCT-CC 27225
DB 731 AAGTACCTGGACTACGAGCGGCGACCAACGCGTGTCTAATTAATTAATTTTGTGT 790
DB 27224 TGTGAGCTGGGAATACAGGCGGCGACCAACGCGGCTAATTTTATA--TTTCAAGT 27167
OY 791 AGAGACTGGGCTTACTAGTGTGCGCAGGCTGTGTTAACTCTGCTGCTTCAAGCAATCC 850
DB 27166 AGAGTCGACAGTTTCACTATTTTCCAGGCTGTGTCTCAAACTCCGACCTCAAGTATCC 27107
OY 851 TCTTACCTTGGCATCCAAAGTCTGAGATTAACAGGCTGAGCCCAATGTGCGCTACT 910
DB 27106 ACCCGCGAGGCTCCCAACGCTGAGGATTAATAGGCTGAGCCCACTGACCTGCTTT 27047
OY 911 TATTTCTTACATTCATCTTTCCAAATGTAATGATCCACAAACAGGATTAAGTGC 970
DB 27046 CTGTACTCTTATTTCTATTTTCACTGCTGTTCTTTTGGTTTCAATATGATTAATTT 26987
```


QY	971	CTATTTTCCTCCCTCTCTTTTGTGAGACAGAGTCTCACTTCATCACTCAACCTCCGGTCA	1030
Db	2696	TGTTCTGTAGCTTAAATAAT--ATATATTTTAAAGCTGGAAAGCAGTGCATGATCTCA	26929
QY	1031	GCTCACTGCACACTCTGCTCTCCGGGTTCAAGATTTCTCTGCTAGCCTTGAAGTA	1090
Db	26928	GCTCACTGCACAACCTCCACCTCCAGAGGCTCAAAATTTATCTGTGACTCACTCTCTGAAGTA	26869
QY	1091	GCTGGAAATTCAACCGTGCACACACAGCTTGGCTAATTTTGTATTTTGTAGCAAGAT	1150
Db	26868	GCTGGAAATTCAAGCAGACGACACACAGTCCCACTAA--TTTTGTATTTTTCAGTGAAGG	26810
QY	1151	GGGGTTTTTACCATTGTCGCCAGGCTGGTCTCAAACTCTGAACCTCAAGGATCTGCTGAC	1210
Db	26809	GAAGTTTCACCATGTTTCTTAAAGGTGATCTTAAATCTCTGACCTGAGAGATCCGCCGCG	26750
QY	1211	CTCACTCTCCCAAAAGTCTGGAATTTATAGCGGTGAGTCACTGTGCTGCGGCATTACTGT	1270
Db	26749	GTCAGCCTCTCCAAAGTCTGGGATTAACAGGCATTAAGCTGTGTCGACCTCTTCTGAT	26690
QY	1271	CTATTTTCTTATTTGCTATATGCCAGATCTTAAAGCAGTGTGCAATATAGAGGTGCT	1330
Db	26689	TCTTTATTTCTGTTTCAATGATCTGTCTTTTGGTTTCAAGTATCAATATTTTGTTTT	26630
QY	1331	CAATTAATTAATGTAGTGAATGCACAGCCTAGATATAAACTTTCTTTTCTTTTAAAAAC	1390
Db	26629	GTTTCTATAGCTTAAAAATATATGATTTT---TTTGAGATAGGGGTCTCACTGTGACACC	26573
QY	1391	AATTTTGACAATTGGACAAATTAATACAATCTTGCATCTGTGTTTCTACTTATCACTT	1450
Db	26572	AGGCTGAGGTGACTGTGGTGTGATCAAGGCTCACTGATCTCTCAACCTCTTAATACAA--T	26514
QY	1451	TGTTATGACTTTTTCATATTTGCTCAAACTCTTATTTGTAAGTGTTTTTCATTTGTAACTA	1510
Db	26513	TGATCTCTCCACCTCAACCTCCAAAGTAGTGCACACATGTTTGGCTAATTTTAAAAT	26454
QY	1511	TTTTAGTCACTGAATTAATATAGCTTAAATTTTCTTATTAATCTCTGCTCCAATTAGAA	1570
Db	26453	TTTTTGTAGAAACAGAGTCTTGGCATGTCAACCAAGGCTGGTCTTGAATCTCGGGCTGAA	26394
QY	1571	GGCCAAATTTTACAATCTGATGAAAGGTATGAAACCTCTCCCAAGAAATTCACACACA	1630
Db	26393	G---AGATCCGCTGACTCACTCTCCCAAGTCTGGAATTAAGGTGTGCACCACTTGG	26337
QY	1631	CACACACACTCACACAGTTTTTTTTTATATGTTTGCAACTAAGCAGAA-----A	1682
Db	26336	CCGAGCCAAAAATATGTTTTTTGGCATGCTCAAGAGTGCGGAGTTCAAGACAGCCTGG	26277
QY	1683	CTGTCAATTAAAGGATGTTTGTTCATATTAATTAATAATCACTAGTTGGGCACTGACT	1742
Db	26276	CCAACATAGGCAAACTGTCTCTTAAAGAAATACAAAAATTAATCTGGGCGACAGTGGCT	26217
QY	1743	CAAGCCTGTAAACCAAGTACTTTTGGAAAGTCAAGGTGGGGTCACTTGAGGTGAGAG	1802
Db	26216	CATGTCTGTAAATTCAGTACTCTGGAAAGGCGAAGGAGGTGATCACTGAGGTCAAGAG	26157
QY	1803	TTTGAGACCAAGCCTGTGCAATATGTGTAAACCTTATCTTACTTAAATAATCAAAAAATTG	1862
Db	26156	TTTAAAGACCAAGCCTGGGCAACCTGTGTAAACCTGTCTTACTTAAATAATCAAAAAATTG	26097
QY	1863	CTGGGTGTATGATGATGCTGTGATAGTCCAGCTACTCTGGGAGGCTGAGGCAAGGAATT	1922
Db	26096	CTGGGCGGTGTGACCGGTGCTCTTAATTCCTCAGTACTCAAGAGGCTGAGGCGAGGAATTC	26037
QY	1923	GCTTGAACCTTGGAGGCGAGAGTTTGGACGTGAGCCGAGATCCACACACTGCACTCCAGCT	1982
Db	26036	ACTTGAACCTTGGAGGTGAGAGTTTGGACGTGAGCCGAGATCAATGCCATTGCACTTAGCT	25977
QY	1983	GGGGGACACAGCAGATCTTATCTCAAAAAAATAAATAATTAATAATTA 2030	
Db	25976	GGGGGACACAGCAGATCTTGTCTCAAAAAAGAAAAAAGAAAAAAA 25929	

ACN44170/C	12	RESULT
ID ACN44170	standard; DNA; 196686 BP.	
AC	ACN44170;	
XX		
XX	18-NOV-2004 (first entry)	
DT		
XX	Human genomic sequence hCG39530.	
DE		
XX	Cytostatic; carcinoma; lymphoma; cancer; human; gene; ss.	
KW		
XX	Homo sapiens.	
OS		
XX	WO2003073826-A2.	
PN		
XX	12-SEP-2003.	
PD		
XX	28-FEB-2003; 2003WO-US006235.	
PF		
XX	01-MAR-2002; 2002US-00087192.	
PR		
XX	(SAGR-) SAGRES DISCOVERY.	
PA		
XX	Morris DW;	
PI		
XX	WPI; 2003-328604/31.	
DR		
XX		
PT	Recombinant nucleic acid useful for diagnosis and treatment of carcinoma	
XX	comprises a nucleotide sequence.	
XX	Claim 1; SEQ ID NO 484; Opp; English.	
XX		
XX	The present invention relates to novel DNA and protein sequences which	
XX	are associated with carcinomas. The sequences are useful for: (i) for	
CC	screening drug candidates; (ii) for screening of bioactive agent capable	
CC	of binding to Carcinoma Associated Protein (CAP); (iii) for screening of	
CC	a bioactive agent capable of modulating the activity of CAP; (iv) for	
CC	evaluating the effect of a candidate carcinoma drug; (v) for diagnosing	
CC	carcinoma; (vi) for inhibiting the activity of CAP; (vii) for treating	
CC	carcinoma; (viii) for neutralizing the effect of CAP; (ix) as a biochip;	
CC	(x) for diagnosing carcinoma or a propensity to carcinoma; and (xi) for	
CC	determining Carcinoma Associated (CA) gene copy number. In addition, the	
CC	CA genes are useful as DNA vaccines and the CAP are useful as markers of	
CC	carcinoma including lymphoma. The present sequence is one such CA coding	
CC	sequence. Note: This patent is an equivalent to basic patent	
CC	US200218286A1, for which no sequence data was published	
CC		
XX		
XX	Sequence 196686 BP; 53978 A; 42758 C; 43862 G; 55372 T; 0 U; 716 Other;	
XX		
XX	Query Match 16.6%; Score 414.2; DB 11; Length 196686;	
XX	Best Local Similarity 60.7%; Pred. No. 3.7e-75;	
XX	Matches 869; Conservative 1; Mismatches 464; Indels 97; Gaps 8;	
XX		
XX	605 TTTTGTGTTGTTGTTTGAAGACAGGGCTTCTCTGTCAACCCAGGCATGACAGTG 664	
XX	TTTCATCATTTATTCATTCAGATGGGGTCTTCTCTGTGTCACGGCTGAGTGAATG 98934	
XX		
XX	665 GTGGAACATAGGCACTGTGAGGCTCAACCTCCTGAGCTCAAGGATTCGCTGACCTGAG 724	
XX	GTGCAATCTCGGGCTCATGTGCAACCTCCACCTCCAGGATTCMAATGATTCCTCGCTCCAG 98874	
XX		
XX	725 CCTCCCAAGTAGCTGGAGCTACGAGCGGTGACACCAACGCGCTGGCTTAATTAATAAATTT 784	
XX	CTCTCTGAGTATCTGGGATTTACAGACACCTGCGCACACACCCGGCTAATTTTGTGTA--TT 98816	
XX		
XX	785 TTTTGTGAGAGACTGGGCTTACTACTGCTGGCCAGGCTTGTCTTAACTCTGGCTTCAAG 844	
XX	TTTGTGTAATGATGGAGTTTCAACATGTGTGGCCAGCTGTGCTCAAACTCTGACCTCAAA 98756	
XX		
XX	845 CAATCCCTCTACCTGTGGCAATCCCAAAAGTGTGGGATTTAAGAGGGGTGAGCAACCATGTGG 904	
XX	GGATTCACCAACCTGTGGCTCCCAAAAGTGTGGGATTTAAGAGGATGAGCCTACGTGGCCCA 98696	

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QY 905 GCTACTATTCTTTACATTCATCTTTCGAATAGTAAGATCCACAGACGGAT 964
Db 98695 GCCATGTACATTATTAATTTACTATATCTCTTTGGT-----CACCATCTTGTCTT 98644
QY 965 TACTGCATATTTCTCTCTCTTTTGTGAGACAGAGTCTCATCTTCACCTCAACCTC 1024
Db 98643 TTTTCTTTTCTTTTCTTTTCTTTTGTGAGACAGAGTCTGTCTGTGCGCCAGGCTGG 98584
QY 1025 CGT-----TCAGCTCATGCAACCTCTGCTCTCCGGGTTCAAGTATCTC 1070
Db 98583 AGTGCAGTGGCGGCATCTCGGCTCATGCAAGCTCCGCTCCCTGGCTCACGCCATCTCTC 98524
QY 1071 CTGCTTAAGCCTCTGAGTACTGGAATTAAGCGGTGCACACACATGCTGGCTAATTT 1130
Db 98523 CTGCTCATGCTCTCAAGATAGCTGGACTACATGTGCCACACACACCGCCGAGAAATTT 98464
QY 1131 TTTGTATTTTATGACAGATGGGTTTATACATGTGGCCAGGCTGTCTCAAACTCTCG 1190
Db 98463 TTTGTATTTTATGAGAGACGGGGTTTCACGGTGTACCGAGATGTCTCGATCTCTG 98404
QY 1191 ACCTCAAGTATCTGCTGCTCTCACTCTCCAAAGTCTGGAATTAATGCGTGAATC 1250
Db 98403 ACCTC--GTGATCCACCCACCTCAAGCTCTCCAAAGTCTGGAATTAAGAGTAC 98346
QY 1251 TGTGCTGGCCGATTAAGTCTGATTTCTTATTTCTATATCCCGAGTCTAGAGCAGTG 1310
Db 98345 TGGGCGCGGCCACCACTGCTCTCTGATTTCTTATTTTCTTCTTGAGAG----- 98291
QY 1311 TCTGACATATAGTAGTGTCTCAATTAATTAATGATGAATGCAAGCTAGATATACTT 1370
Db 98290 -----AGGAATTC 98283
QY 1371 TCTTTTCTTTTAAACAATCTTGACAACTTGCAGAAATATACATCTTGATTC 1430
Db 98282 TATATGTTCTTCAACCATATCTCTAATATACCTTCTTCTCTAATATCTAATACTA 98223
QY 1431 TGTCTTTTCACTTACCTGTGTATGACCTTTTCATATGCGCTCAAACTTATGTTA 1490
Db 98222 TATATACATACATATAGTGTATATATATATATGATATATATATATATATAT 98163
QY 1491 CTGTTTTTCACTTGTACTATTTTATGTCACGTGAATATATGCTTAATTTGCTTATCAT 1550
Db 98162 ATATATATACACAGTCTTATTTATGTTATTCATTTGCTTAATACATATGTTTTCAT 98104
QY 1551 CCTCTGCTCACTTTAGAGGCCAAATTTACAAATCTGATGAAGCTATGAACTCTC 1610
Db 98103 -----AAACCTATTAAGAACCTTTAAACAAGTATGAGTGAAGAACCTTGAAGTTC 98051
QY 1611 CCCAGAGAAATACACACACACACACACACACAGTCTTTTATTTTATGTTTGCAC 1670
Db 98050 CCAGATTAATTAATTTGTCTCAAAAGTCTCATCTTTGTTTATGTTTATCTCTCAAT 97991
QY 1671 TAAAGCAAGAAACCTGCATTAGAGATGTTTGTCTATATTAATTAATAAATA----CTCA 1726
Db 97990 TATTTATCTATATGGCAATTAACAATATTTTGTGTTGTCACAAATAATAGTCTCTG 97931
QY 1727 GTTGGCACAGTGCATCAAGCCTGTAAACACAGTACTTTGGAAGTCCAAAGTGGGTGAT 1786
Db 97930 GCTGACACAGGTGCTCAGCCTGTAAATCCAGCACTTTGGAGGCGCAGAGGGGTGAT 97871
QY 1787 CACTGAGGTGAGAGTGTGAGACACGCTGTGCAATATGTAAGAAACCTATATCTACTA 1846
Db 97870 CAC--CAGGTACAGAGATGGAACATCTCTGGCCAAACATGTAATCTCTCTCTACTA 97813
QY 1847 AAAATATCAAAAATTTAGCTGGGTGTAGTATGATGCAATGCTTGTCTCTGAGAG 1906
Db 97812 CAATATCAAAAATTTAGCTGGGTGTAGTATGCAATGCGCTGTAGTCTCAAGTACTCTG 97753
QY 1907 CTGAGGCAAGAAATTTGTTGAACCTGGAGGCAAGGTTTGCATGAGCCGAGATCTCCAC 1966
Db 97752 CTGAGGCAAGAAATTTGTTGAACCTGGAGGCAAGGTTTGCATGAGCCGAGTCAAGC 97693
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QY 1967 CACTGCACTCCAGCCTGGCGACACAGCAGAGACTATCTCAAAAATAA 2017
Db 97692 CACTGCACTCCAGCCTGGGTGACAGAGAGACTATCTCAAAAATAA 97642

RESULT 13
ADP68568/c
ID ADP68568 standard; cDNA; 86000 BP.
XX
AC ADP68568;
XX
DT 09-SEP-2004 (first entry)
XX
DE Human PPAR-alpha cDNA.
XX
KM cytosolic; gene therapy; PPAR-alpha;
KM peroxisome proliferator-activated receptor-alpha; PPAR-alpha modulator;
KM PPAR-alpha associated disorder; hyperproliferative disorder; human; gene;
86.
XX
OS Homo sapiens.
XX
EN US2004115637-A1.
XX
PD 17-JUN-2004.
XX
PE 11-DEC-2002; 2002US-00317500.
XX
PR 11-DEC-2002; 2002US-00317500.
XX
PA (ISIS-) ISIS PHARM INC.
XX
PI McKay R, Dobie KM;
XX
WP1; 2004-449378/42.
XX
DR
XX
PT New oligonucleotide compound that inhibits expression of PPAR-alpha,
PT useful for preparing a composition for treating hyperproliferative
PT disorders, e.g. cancer.
XX
PS Claim 1; SEQ ID NO 4; 121bp; English.
XX
CC The invention describes a compound, having a sequence comprising 8-80 bp
CC targeted to a nucleic acid encoding PPAR-alpha (peroxisome proliferator-
CC activated receptor-alpha), that specifically hybridizes with the nucleic
CC acid encoding PPAR-alpha comprising 86001-bp sequence and inhibits
CC expression of PPAR-alpha. Also described are: a method of inhibiting the
CC modulator of PPAR-alpha; a diagnostic method for identifying a disease
CC state; a kit or assay device comprising the compound; and a method of
CC treating an animal having a disease or condition associated with PPAR-
CC alpha. The oligonucleotide compound is useful for preparing a composition
CC for treating hyperproliferative disorder e.g. cancer. This sequence
CC represents a human peroxisome proliferator-activated receptor-alpha (PPAR
CC -alpha) cDNA.
XX
SQ Sequence 86000 BP; 22822 A; 19623 C; 20377 G; 23178 T; 0 U; 0 Other;

Query Match 16.6%; Score 413.6; DB 12; Length 86000;
Best Local Similarity 61.0%; Pred. No. 4.1e-75;
Matches 878; Conservative 1; Mismatches 490; Indels 71; Gaps 10;

QY 605 TTTTGTGTGTGTATGAGACAGGGTCTGTCTGACCCAGGAGTGAAGCAGATG 664
Db 69228 TTTTGTGTGTGTATGAGACAGGGTCTGTCTGACCCAGGAGTGAAGCAGATG 69169
QY 665 GTGCAACATAGTCACTGACGCTCAACCTCTGAGCTCAAGGATCTGCTGACCTGAG 724
Db 69168 GTGCAATCTCACTGACGCAACCTCCGCTCCGAGTTCAAGCTATCTCTGCTCAT 69109
QY 725 CTCTCCAGTACTGGGACTAGAGCGTGCACACACAGCCGCTGATTAATAAATTT 784
Db 69108 CTCTCCAGTACTGGGACTAGAGCGTGCACACACAGCCGCTGATTAATAAATTTTGTAA--TT 69051
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QY 785 TTTTGTAGAGACTGGGTCTTACTAGTGGCGAGGCTTGTCTTAACTCTGGCTTGAAG 844
D 69050 TTTTATAGAGACGGGGTTTGGCATTTTGGCATTTGGCATTTGGCATTTGGCATTTG 68991
QY 845 CAATCTCTTCACTTGGGATCCCAAGTGTGGGATTAAGAGGGTGAACCAATGTGCG 904
D 68990 TGAATCCACCCGCTCGGCTTCCAAAGTGTGGGATTAAGAGTGAACCAACCGACCTG 68932
QY 905 GCTACTTATTTCTTACATTCATCTTCCAAATAGATAGATCCACAGAACGGGAT 964
D 68931 GGAGACTTTTATTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCT 68872
QY 965 TACTGCTATTTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCT 1024
D 68871 CTCCTCTGTGCGCCAGGCTGAGTGCATGGCCGATC----- 68833
QY 1025 GCTTCACTCACTGCAACTCTGCTCCGGGTTCAGAGATTCCTCTGCTTAAAGCTCC 1084
D 68832 ---TCAGTCACTGCAACTCTGCTCCGGGTTCAGAGATTCCTCTGCTTAAAGCTCC 68776
QY 1085 TGAATAGCTGAATTAACAGGCTGCAACCACTGCTTAAATTTTGTATTTTATG 1144
D 68775 CGAGTAGCTGGAGCTACAGGATGTGCCACCAACCCAGCTTA-TTTTGCATTTCTAGT 68717
QY 1145 AGAGATGGGGTTTACCATGTTGCCAGGCTGTGTCTGAACTCTGACCTGAGTATCT 1204
D 68716 AGAGACGGGGTTTACCATGTTGCCAGGCTGTGTCTGAACTCTGACCTGAGTATCT 68657
QY 1205 GCTGCTCACTGCTTCCCAAGTGTGAAATTAAGGCTGAGTCACTGCTGCTGGCGAT 1264
D 68656 ACCCACTCACTGCTTCCCAAGTGTGAAATTAAGGCTGAGTCACTGCTGCTGGCGA- 68598
QY 1265 TACTGTCTATTTCTTATTTCTTATTTCTTATTTCTTATTTCTTATTTCTTATTT 1324
D 68597 ---CTTTATTTCTTATTTCTTATTTCTTATTTCTTATTTCTTATTTCTTATTTCT 68542
QY 1325 GGTGCTCAATTAATATGATGATGATGATGATGATGATGATGATGATGATGATGAT 1384
D 68541 GCAATTCATCATATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 68489
QY 1385 TAAACAATCTTGAACAATTTGCAAGATTAATTAATTAATTAATTAATTAATTAAT 1444
D 68488 -GGAACGAATCTGGAATTAATTAATTAATTAATTAATTAATTAATTAATTAATTAAT 68436
QY 1445 TCACCTGTTATGACTTTTCTTATTTGCTTCAAACTTTATTTGTTTCTTCTTCT 1504
D 68435 CCGAC-AGAAAGGCACTTGTAAATCGACATCTGCACTTGTCTTCTTCTTCTTCTG 68377
QY 1505 TTACTATTTTGTGACTGAATTAATTAATTAATTAATTAATTAATTAATTAATTAAT 1564
D 68376 ---GATCTTGACCTGCGGTGCACTTGTATTAACAGCTTGTGATTAATTTGCTG 68321
QY 1565 TTAAAGGCGCAATTTTCAAAATCTGATGAAGTATGAACCTCTCCACAGAGAAATCA 1624
D 68320 GAAAGAAAGCCCTGTAGAGGTAAATCAAGTGAATTTAAAGACCTGGGCACTGAGAAAC 68261
QY 1625 CACACACACACACTACACACAGTTTCTTAAATTTTGAACCTGAGCAAGAAACC 1684
D 68260 AGCCCACTTGTCTTACTACACTTTTGTGTTGCTTATTTGTTCTTCTTCTTCTTCT 68201
QY 1685 TGCATTAAGAGATTTTGTCTTATTAATTAATTAATTAATTAATTAATTAATTAAT 1744
D 68200 AATTTTGAAGAAAGTATTC-----CAGGCCAGGCACTGTGATCA 68160
QY 1745 AGCTGTAAACCACTTCTTGAAGTCAAGTGTGATCACTTGAAGGTGAAGTT 1804
D 68159 TGCTGTATTCAGCACTTTAGAGGTGAGGGGGGTGATCACTGAGGTGAGGAT 68100
QY 1805 CGAATCCAGCTGTGATTAATTTGAAACCTTATCTTACTTAAATTAATTAATTAAT 1864
D 68099 CGAATCCAGCTGTGATTAATTTGAAACCTTATCTTACTTAAATTAATTAATTAAT 68040

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QY 1865 GGTGTAGTATGATGATCTCTGATCTCCAGTACTCGGAGGCTGAGGCAAGAAATTGC 1924
D 68039 GGGATGTATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 67980
QY 1925 TTGAACCTGGAGGCGAGAGGTTGCACTGAGCCGAGATCCCACTGCACTCCAGCTGG 1984
D 67979 TTGAACCCAGAGGCGAGAGGTTGCACTGAGCCGAGATGATGATGATGATGATGAT 67920
QY 1985 GCGACAGAGGAGCTCTATCTCAAAAATTAATTAATTAATTAATTAATTAATTAAT 2044
D 67919 GCGACAGAGGAGCTCTATCTCAAAAATTAATTAATTAATTAATTAATTAATTAAT 67860

RESULT 14
AD097523/c
ID AD097523 standard; DNA; 215974 BP.
XX
XX AD097523;
AC AD097523;
DT 07-OCT-2004 (first entry)
XX
DE Human cancer associated sequence HD09-008, SEQ ID 500.
DE
DE Cytostatic; Gene Therapy; cancer; leukemia; lymphoma; Human; de.
XX
XX Homo sapiens.
XX
XX W02040460304-A2.
XX
XX 22-JUL-2004.
XX
XX 22-DEC-2003; 2003WC-US041389.
XX
XX 27-DEC-2002; 2002US-00330773.
XX
XX (SAGR-) SAGRES DISCOVERY INC.
XX
XX Morris DW, Malandro MS;
XX
XX WPI; 2004-543781/52.
XX
XX New isolated cancer associated nucleic acids comprising at least 10
XX contiguous nucleotides, useful for diagnosing, preventing and/or treating
XX cancers such as leukemia and lymphoma.
XX
XX Claim 1; SEQ ID NO 500; 199bp; English.
XX
XX The present invention relates to cancer associated sequences (AD097025-
XX AD098004). The sequences are useful for the diagnosis, prevention and/or
XX treatment of cancer, such as leukemia and lymphoma. Note: The sequence
XX data for this patent did not form part of the printed specification, but
XX was obtained in electronic format directly from WIPO at
XX ftp.wipo.int/pub/published_pat_sequences.
XX
XX Sequence 215974 BP; 56601 A; 42897 C; 46516 G; 65586 T; 0 U; 4374 Other;

Query Match 16.6%; Score 413.6; DB 12; Length 215974;
Best Local Similarity 59.6%; Pred. No. 5e-75;
Matches 847; Conservative 0; Mismatches 529; Indels 46; Gaps 7;

QY 610 TTTTGTGTTTGAAGACAGAGGCTGTGCTCTGTCACCCAGGATGAGCAGTGTGCA 669
D 4846 TGTCTTTGTTTGAAGACAGAGGCTGTGCTCTGTCACCCAGGATGAGTACAGTGCACA 4787
QY 670 ACCATAGTCACTGACCTCAACTCTGAGCTCAAGGATGCTGACCTCAGCCTCC 729
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GenCore version 5.1.7
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OM nucleic - nucleic search, using sw model

Run on: February 17, 2006, 09:56:20 ; Search time 12639 Seconds
(without alignments)
11248.158 Million cell updates/sec

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Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 5883141 seqs, 28421725653 residues

Total number of hits satisfying chosen parameters: 11766282

Minimum DB seq length: 0

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Post-processing: Minimum Match 0%
Maximum Match 100%

Listing first 45 summaries

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4: gb_om:*
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11: gb_by:*
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13: gb_vl:*
14: gb_hcg:*
15: gb_pl:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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ALIGNMENTS

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DEFINITION Homo sapiens phospholipase A2, group 1B (pancreas) (PLA2G1B) gene, complete cds.
ACCESSION AY438977
VERSION AY438977.1 GI:37953284
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Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Homidae; Homo.
REFERENCE 1 (bases 1 to 8368)
Rieder,M.J., Livingston,R.J., Daniels,M.R., Chung,M.-W., Miyamoto,K.E., Nguyen,C.P., Nguyen,D.A., Poel,C.L., Robertson,P.D., Schackwitz,W.S., Sherwood,J.K., Witrak,L.A. and Nickerson,D.A.
Direct Submission
JOURNAL Submitted (16-OCT-2003) Genome Sciences, University of Washington, 1705 NE Pacific, Seattle, WA 98195, USA
COMMENT To cite this work please use: NIHES-SNPs, Environmental Genome Project, NIHES ES15478, Department of Genome Sciences, Seattle, WA (URL: <http://lepp.gs.washington.edu>).

FEATURES
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DEFINITION	Sequence 1 from Patent WO02122562.		
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VERSION	AX377239.1	GI:19573528	
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SOURCE			
ORGANISM	Homo sapiens (human)		
	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;		
	Mammalia; Euteria; Euarchontoglires; Primates; Catarrhini;		
	Homnidae; Homo.		
REFERENCE			
AUTHORS	Kazemi, A., Kitem, S.E. and Koshy, B.		
TITLE	Haplotypes of the pla2g1b gene		
JOURNAL	Patent: WO 0212562-A 1 14-FEB-2002;		
	Genaisance Pharmaceuticals, Inc. (US)		
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variation	9531	/note="PSS: polymorphic base G or A"
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QY	1	GTCTGTCTACCTGCTGTCCAGCTGGGTAAACAGACCACTCTGTCTCAAAAAAAAAAATG 60
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QY	61	CTTTCAATTAATATATGATTAAGAGCTTAATATTTTCAAGCATTAGATCAATTTCTCC 120
DB	6576	CTTTCAATTAATATATGATTAAGAGCTTAATATTTTCAAGCATTAGATCAATTTCTCC 6635
QY	121	TGAAGCATCTTGGGGAATCAATCCCACTGTCTGAGAGTGGGAGGTGAGGGCTGAC 180
DB	6636	TGAAGCATCTTGGGGAATCAATCCCACTGTCTGAGAGTGGGAGGTGAGGGCTGAC 6695
QY	181	CTATTGCTCTGCACTTACTCTATCTCAAGCTGTCCCTCCACTTTTCAAGTGTGCTCCAGA 240
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DB	6756	CACATGAACAATGCTAAGCAAGGCAAGAGCTGGAAGCTGTAAATTTCTGCTGACA 6815
QY	301	MMCCCTACACCAACCACTTATCATCTCTGTCTGTGCTGGCAATCACTGTAGCAGTA 360
DB	6816	ACCCTACACCAACCACTTATCATCTCTGTCTGTGCTGGCAATCACTGTAGCAGTA 6875
QY	361	GATTATCCCTTCTTGAAGCTATGAATCTTAGTGTCTCTCAAGTGGCCGGGGGGAATA 420
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QY	421	ATAGTAACAACAAGCAGATTAATTAAGTTAAATTTCTGTGGGAGTGTCTCTTA 480
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DB	7356	CAAGCAATCTCTCACTTGGCATCCCAAGTGTGGGATTAACAGGGGTGAGCCACATG 7415
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RESULT 3
 AC003982/c 122302 bp DNA linear PRI 18-MAR-1999
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 DEFINITION Homo sapiens PAC clone 166H1 from 12q, complete sequence.
 AC003982
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 SOURCE
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 Homo sapiens
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 Homidae; Homo.
 1 (bases 1 to 122302)
 Sultston, J.E. and Waterston, R.
 Toward a complete human genome sequence
 Genome Res. 8 (11), 1097-1108 (1998)
 9847074
 2 (bases 1 to 122302)
 Bradshaw, H., Wu, X. and Ozersky, P.
 The sequence of Homo sapiens PAC clone 166H1
 Unpublished (1999)
 3 (bases 1 to 122302)
 Waterston, R.
 Direct Submission
 Submitted (13-JUN-1998) Department of Genetics, Washington
 University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
 4 (bases 1 to 122302)
 Waterston, R.
 Direct Submission
 Submitted (18-MAR-1999) Department of Genetics, Washington
 University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
 SUBMITTED BY: WUSC
 Genome Sequencing Center
 Department of Genetics
 Washington University
 St. Louis MO 63108, USA
 http://genome.wustl.edu/gsc
 mailto:sapiens@watson.wustl.edu

NOTICE: This sequence may not represent the entire insert of this
 clone. It may be shorter because we only sequence overlapping
 clone sections once, or longer because we provide a small overlap

between neighboring data submissions.

This sequence was finished as follows unless otherwise noted:
all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:

This clone was originally isolated in the laboratory of Professor Greame Bell, Howard Hughes Medical Institute and Departments of Biochemistry and Molecular Biology, and Medicine, The University of Chicago, Chicago, IL, USA. The clone was provided by the Laboratory of Dr. Roger Cox at The Wellcome Trust Centre For Human Genetics, Oxford, UK. Some contig information was also obtained from Yamagata et al., Nature 384:455-8 (1996).

SOURCE INFORMATION:

This clone was derived from human PAC library RPCT-1, prepared by Pieter de Jong and coworkers at the Roswell Park Cancer Institute (<http://bacpac.med.buffalo.edu>) using the method described by Ioannou et al., Nature Genetics 6:84-9 (1994). The library is from one male donor.

The clone may be obtained either from Genome Systems, Inc.

(<http://www.genomesystems.com>) or Research Genetics, Inc.

(<http://www.regen.com>); or from Pieter de Jong.

VECTOR: pCYPAC2

NEIGHBORING SEQUENCE INFORMATION:

The clone sequenced to the left is 278C19; the clone sequenced to the right is 15E1. Actual start of this clone is at base position 1 of 166H; actual end is at 122302 of 166H.

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Query Match 99.9%; Score 2497.4; DB 8; Length 122302;

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 Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
 Homnidae; Homo.
 1 (bases 1 to 220384)

REFERENCE
 AUTHORS
 Muzny, D.M., Adams, C., Adio-Oduola, B., Ali-osman, F.R., Allen, C.,
 Alstrooks, S.L., Amaral, H.C., Are, J.R., Ayala, M., Banks, T.,
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TITLE
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Homi, F., Howard, S., Huber, J., Hulyk, S., Hume, J., Jackson, L.E.,
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 Direct Submission
 Unpublished
 2 (bases 1 to 220384)
 Worley, K.C.
 Direct Submission
 Submitted (11-AUG-2000) Human Genome Sequencing Center, Department
 of Molecular and Human Genetics, Baylor College of Medicine, One
 Baylor Plaza, Houston, TX 77030, USA
 3 (bases 1 to 220384)
 Worley, K.C.
 Direct Submission
 Submitted (26-MAR-2002) Human Genome Sequencing Center, Department
 of Molecular and Human Genetics, Baylor College of Medicine, One
 Baylor Plaza, Houston, TX 77030, USA
 On Mar 26, 2002 this sequence version replaced gi:18449664.
 ----- Genome Center
 Center: Baylor College of Medicine
 Center code: BCM
 Web site: http://www.hgsc.bcm.tmc.edu/
 Contact: hgsc-help@bcm.tmc.edu
 ----- Project Information
 Center project name: HGM
 Center clone name: RP11-836M1
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 Sequencing vector: Plasmid;
 Sequencing vector: M13;
 Chemistry: Dye-terminator Big Dye; 53% of reads
 Chemistry: Dye-terminator Big Dye; 53% of reads
 Assembly program: Phrap; version 0.990329
 Consensus quality: 234783 bases at least Q40
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 Consensus quality: 245671 bases at least Q20
 Estimated insert size: 219187, sum-of-coverage estimation
 Quality coverage: 8.9x in Q20 bases; sum-of-coverage estimation

 NOTE: Estimated insert size may differ from sequence length
 (see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html).
 NOTE: This is a "working draft" sequence. It currently
 consists of 11 contigs. The true order of the pieces
 is not known and their order in this sequence record is
 arbitrary. Gaps between the contigs are represented as
 runs of N, but the exact sizes of the gaps are unknown.
 This record will be updated with the finished sequence
 as soon as it is available and the accession number will
 be replaced.
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Center clone name: RP11-144B2
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Sequencing vector: Plasmid; M77789
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Assembly program: Phrap; version 0.990329
Consensus quality: 191640 bases at least Q40
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Quality coverage: 0x in Q20 bases; agarose-fp estimation
Quality coverage: 11.7x in Q20 bases; sum-of-contigs estimation
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* (see http://www.hgsc.bcm.tmc.edu/docs/genbank_drafc_data.html).
* NOTE: This is a 'working draft' sequence. It currently
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* is not known and their order in this sequence record is
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* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
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 Kaul, R.K., Olson, M.V., Zhou, Y., James, R.A., Rouse, G., Wu, Z.,
 Saenphimachak, C., Phelps, K.A., Buckley, D., Kibukawa, M., Raymond, C.
 and Haugen, E.D.
 Direct Submission
 Unpublished
 2 (bases 1 to 190162)
 Bubb, K.L., Desmarais, C.L., Ramsey, S.A. and Hubley, R.M.
 Submitted (07-AUG-1999) Human Genome Center, University of
 Washington, Box 352145, Seattle, WA 98195, USA
 3 (bases 1 to 190162)
 Kaul, R.K., Zhou, Y., James, R.A., Raymond, C., Haugen, E.D. and
 Olson, M.V.
 Direct Submission
 Submitted (30-OCT-2000) Genome Center, University of Washington,
 Box 352145, Seattle, WA 98195, USA
 4 (bases 1 to 190162)
 Kaul, R.K., Olson, M.V., Zhou, Y., James, R.A., Rouse, G., Wu, Z.,
 Saenphimachak, C., Phelps, K.A., Buckley, D., Kibukawa, M., Raymond, C.
 and Haugen, E.D.
 Direct Submission
 Submitted (26-MAY-2002) Genome Center, University of Washington,
 Box 352145, Seattle, WA 98195, USA
 On May 26, 2002 this sequence version replaced gi:11038533.
 ----- Genome Center
 Center: University of Washington Genome Center
 Center Code: UWGC
 Web site: http://www.genome.washington.edu
 Contact: uwgchgs@u.washington.edu
 ----- Project Information
 Center project name: chr-7
 Center clone name: RP11-305M3 (d3s195)
 ----- Summary Statistics
 Assembly program: Phrap: version 0.990319
 Consensus quality: 189869 bases at least Q40
 Consensus quality: 190110 bases at least Q30
 Consensus quality: 190161 bases at least Q20
 Insert size: 190162; sum-of-contigs
 Quality coverage: 9.7x in Q20 bases; sum-of-contigs

 Overlapping Sequences:
 5': RP11-382M23 (UWGC:d3s734g) AC093149 1340-bp overlap
 3': RP11-448A19 (UWGC:d3s705) AC078846 8765-bp overlap

 Sequence Quality Assessment:
 This entry has been annotated with sequence quality
 estimates computed by the Phrap assembly program.
 All manually edited bases have been reduced to quality zero.
 Quality levels above 40 are expected to have less than
 1 error in 10,000 bp.
 Base-by-base quality values are not generally visible from the
 GenBank flat file format but are available as part
 of this entry's ASN.1 file.

This sequence was finished as follows unless otherwise noted:

all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., Phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest.

Sequence Validation:

This sequence has been validated by Multiple Complete Digest fingerprinting. Comparison of the experimentally derived digest fragments with sequence-predicted fragments is given below. The electronically-digested sequence consists of both insert and vector, in order to accurately represent the entire circular BAC. Small fragments below a variable cutoff (approximately 400-800 bp) are not resolved in the fingerprint and hence do not appear in the table. There are no significant remaining discrepancies between the experimental and predicted values. Uniquely ordered fragments are separated by dashed lines.

N811

Bg111

ECORI

SeqDerMap	FngPrnt	SeqDerMap	FngPrnt	SeqDerMap	FngPrnt
21251	20929	12417	12174	8696	8794
12148	11918	2067	2041	6	<800
199	<800	5666	5826	1775	1773
3275	3221	254	<800	2000	1991
20379	20929	3818	3836	6313	6324
15555	15150	747	735	3256	3226
675	<800	9196	9237	325	<800
5493	5475	4154	4101	12989	12723
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5810	5831	118	<800	611	<800
9488	9435	2524	2359	1995	1991
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315	<800	26	<800	955	940
3640	3671	4675	4644	2961	2940
428	<800	1801	1803	5264	5205
1144	1119	117	<800	1535	1534
4684	4717	6797	6799	3313	3383
20	<800	3146	3177	3760	3761
492	<800	2376	2359	3061	3110
10935	10879	210	<800	9605	9505
5921	6010	5135	5110	335	<800
529	<800	744	735	2190	2194
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1311	1293	413	<800	13558	13434
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374	<800	296	<800	8080	8019
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1019	1000	466	<800	8756	8794
841	840	927	947	7	<800
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7121	7263	5159	4822		
1122	1119	677	<800		
49	<800	1399	1373		
11928	11918	1565	1554		
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2585	2575	4826	4822		
699	<800	7661	7660		
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		589	<800		

Query Match 18.5%; Score 462.6; DB: 8; Length 190162;
Best Local Similarity 61.3%; Pred. No. 1.3e-83;
Matches 900; Conservative 1; Mismatches 520; Indels 46; Gaps 8;

Qy	604	TTTTTTTGTGTTGTTTGAAGACAGGCTTGTCTGTACACCAAGCATGACAGT	663
Db	31687	TTTTCTTTTAAAGTAGAAGAGCTTGTGCTGACCAAGGCTGGGTGAGT	31746
Qy	664	GGTGCAACATPAGTCACTGACGCTTCACTCTCTGAGTCAAGGATCTGACCTCA	723
Db	31747	AGTGACGTCGTGTTTCACTGACCTTCAATCTCTAGGAACAGGTATCTCCACATTCA	31806

OY	724	GGCTCCAGTGTAGCTGGAGCTAGAGACGTGTACACACACCGCTGGCTAAATTTAAAAATTT	783
Db	31807	GTCTTCTGTAGTACTGGAGCTTAGGCACTTAGGCACCAATGCCACGTAAATTTAAAAAAA	31866
OY	784	TTTTTGTAGAGACTGGGCTTACTACGTGGCGAGCGCTTGTCTTAAATCTCCGTCTTCA	843
Db	31867	TGTTTGAAGACACAGGGTCTTGTCTGTGTTGCTCAGGCTGTGCTTCAACAGCAGCCTTCA	31926
OY	844	GGAATCCTCTTCACTTGGCATGCCAAAGTCTGGGATTTACAGGGGTAGGCCACATGTGC	903
Db	31927	GTGATCTCTGTGCTCATCTCCCAAGTGTGGGATTTACAGGTGTAGGCCACATGTGCC	31986
OY	904	GGCTACTTATTTCTTTACATTCATCTTCCATATAGATGTAGATCCACAGAACAGGGA	963
Db	31987	AGCCCAATTCCTTTTTCTGTTTTTTTTTTAAATTAATTTAAATTTTATTTTATTTT	32046
OY	964	TTACTGCTTATTTTCTTCTCTT-TCCTTTTGAAGACAGAGTCTGACTTCACTCAACC	1022
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OY	1023	TCGGT-----TCAGTCACTGCAACTCTGCTCCCGGGTTCAGATGATTC	1068
Db	32107	GGAGTGCAGTGGCATGATCTCGGCTCACTGAACCTCTGCTCCCAAGTTCAAGATTC	32166
OY	1069	TCCTGCTTAAGCTCTCTGAGTACGTGAATTAACAAGCGTGACACCAATGCTTGGCTAAT	1128
Db	32167	TCCTGCTCAAGCTCTCTGAGTACGTGGATTAACAGTGCCTCCACATATACCGGCTAAT	32225
OY	1129	TTTTTGTATTTTATAGCAGAGATGGGGTTTTTACATGTGGCCAGAGCTGTCTCAACCTCC	1188
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OY	1189	TGACCTCAAGTGTCTGCTGCTCACTGCCAAGTGTGTAATTAAGCGTGAATC	1248
Db	32286	TGACCTCAAGTGTCTGCTGCTGCCCTCCCAAGTGTGGGATTAACAGTGTGAAGCC	32345
OY	1249	ACTGTGCTGGCGGATTACTGTCTAATTTT--CTTATGTGTATATCCCGAATCTAGAG	1305
Db	32346	ACCGAACCCGGCTGCCCAATTTTTTTTTTAAACAAATGTCTTACATCAAGCTAAGAG	32405
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Db	32406	CTGTAAAGGCAATAGACATATGCTTACCTCTAGAATCTTATATTTTGGTGAGATAT	32465
OY	1366	AACCTTCTTTTCTTTTTTAAACAAATCTTGACACTTGGCAGATTAATTAACAATCTTG	1425
Db	32466	ACACATATATTAATATAGATGTCAAGACATATATATATTAAGCCATGACAGATTTATA	32525
OY	1426	CATTTGTCTTTTAC-TTATACCTGTATATGACTTTTATATGTGCTCAACCTTTA	1484
Db	32526	AATCCATATCACTACACAAACAAAGGCTTAGAAGTTCAAGTTCCACTTTGGGAAG	32585
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Db	32586	CTBAGGAGGTGATGACCTGAGGTCAAGAAATTA--AGACAGGTGTGCCAACATGTGTG	32643
OY	1545	ATACATCTCTCTGTCCACTTTAGAGGCCAAATTTACAAATCTGTATGAAGCTATGAAC	1604
Db	32644	AAACCCCTGTCTTTACTAAGATGAACAAATTAAGCTGATGTGTGTGTGTCAGGCTGTAT	32703
OY	1605	CCCTCTCCAGAGAAATACACACACACACACTCACACAGTTTTTTTTTATATGTT	1664
Db	32704	CTTAGCTACTTAGAGGCTAGGACAGAGAAATCGTTGAACCCGAGGCGGAGGCTGCA	32763
OY	1665	TGCAACTAAGACAAAGAACTGCAATTAGAGATGTTG-----1702	
Db	32764	GTGAGCGGAGATCGCGCACTGCACTTACGCTTGGGTGACAGATGAGACTTCATCTCA	32823
OY	1703	TTGATATTTAATTAATAATCACTAGTTGGGACAGTGACTCAAGCTGTAAACACAGTAC	1762
Db	32824	AAAAAAAAAAAAAAAAAGCGCCAGGCGCGGTGTCTCAACTGTGAATCCAGACAC	32883

QY	1763	TTTGGAAAGTCGCAAGGTGGTGGATATCACTTGAAGGTGAGAACTTGACGACCAAGCTGGTCA	1822
Db	32884	TTTGGAGAGCTCGAAGTGGGCGATCA--TAGAGTCAGAACTTGACGACCAAGCTGGCGAA	3294
QY	1823	TATGTGAAACCCCTATCTCTACTATATAAATACAAAATAATTAAGCTGGGTAGTGAATGCATGC	1882
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QY	1883	CTGTATGTCCTCCAGCTACTCTCGGAGGCTGAGGCAGAGAAATTGCTGAACTCGGAGAGAGA	1942
Db	33002	CTGTATGTCCTCCAGCTACTCTCGGAGGCTGAGGCAGAGAAATTGCTGAACTCGGAGAGAGA	3306
QY	1943	GGTTGCAATGAGCGGAGATCCACACTGCACTTCAGGCTGGGCGACACAGCAGACTCT	2002
Db	33062	GGTTGCAATGAGCGGAGATTAATGCTGCACTTCAGGCTGGGCGACAGCAGACTCTC	3312
QY	2003	ATCTCAAAAAAAAAATAATAATAATA 2029	
Db	33122	GTCTCAAAAAAAAAAGAAAAAGAAAA 33148	
RESULT 8	AP000589	123192 bp DNA linear HTG 30-MAY-2000	
LOCUS	AP000589	Homo sapiens chromosome 11 clone CMB9-105NS map 11q13, WORKING	
DEFINITION		DRAFT SEQUENCE, 19 unordered pieces.	
ACCESSION	AP000589		
VERSION	AP000589.3	GI:8118795	
KEYWORDS	HTG; HTGS_PHASE1; HTGS_DRAFT.		
SOURCE	Homo sapiens (human)		
ORGANISM	Homo sapiens		
	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Homiidae; Homo.		
REFERENCE	1 (bases 1 to 123192)		
AUTHORS	Hattori,M., Ishii,K., Toyoda,A., Taylor,T.D., Hong-Seog,P., Fujiyama,A., Yada,T., Totoki,Y., Watanabe,H. and Sakaki,Y.		
TITLE	Homo sapiens 123,192 genomic DNA of 11q13		
JOURNAL	Published Only in Database (1999)		
REFERENCE	2 (bases 1 to 123192)		
AUTHORS	Hattori,M., Ishii,K., Toyoda,A., Taylor,T.D., Hong-Seog,P., Fujiyama,A., Yada,T., Totoki,Y., Watanabe,H. and Sakaki,Y.		
TITLE	Direct Submission		
JOURNAL	Submitted (12-OCT-1999) Masahira Hattori, The Institute of Physical and Chemical Research (RIKEN), Genomic Sciences Center (GSC); Kitasato Univ., 1-15-1 Kitasato, Sagamihara, Kanagawa 228-8555, Japan (E-mail:hattori@gsc.riken.go.jp, URL:http://hgp.gsc.riken.go.jp/, Tel:81-42-778-9923, Fax:81-42-778-9924)		
COMMENT	On May 31, 2000 this sequence version replaced gi:6997479.		
	----- Genome Center		
	Center: RIKEN Genomic Sciences Center (GSC)		
	Center code: RIKEN		
	Web site: http://hgp.gsc.riken.go.jp/		
	Contact: hattori@gsc.riken.go.jp		
	----- Project Information		
	Center project name: HumDrafc11		
	Center clone name: CMB9-105NS		
	----- Summary Statistics		
	Sequencing vector: PCR products; 100% of reads		
	Chemistry: Dye-terminator ET-amersham; 100% of reads		
	Assembly program: Phrap; version 0.990329		
	Consensus quality: 115116 bases at least Q40		
	Consensus quality: 118340 bases at least Q30		
	Consensus quality: 120191 bases at least Q20		
	Insert size: 121392; sum-of-contigs		
	Quality coverage: 5.08x in Q20 bases; sum-of-contigs		

	NOTE: This is a 'working draft' sequence. It currently consists of 19 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs N, but the exact sizes of the gaps are not known.		

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/organism="Homo sapiens"  
/mol_type="genomic DNA"  
/db_xref="taxon:9606"
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Db      63477 ATCATCTTTGATCTTTTTTTTGAAGACAGTCTTGCTGTGTCACACGAGCTGAGTGA 63536
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Db      63537 ATGCGGCATCTCGGCTACACGAACTCTGCTCCGAGGTTACGCAATCTCTGCT 63596
Qy      1077 AAGCTCTGAGTACGCTGGAATTTACAAAGCTGACACCAATGCTGCTGCTAATTTTTT -G 1134
Db      63597 CAGCTCCAGAGTACGCTGGATTAACAGTGGCGGCAACAAGTCTGCTGCTAATTTTTTGT 63656
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RESULT 9
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LOCUS
DEFINITION
Homo sapiens chromosome 15 clone RP11-118016 map 15q15, complete
sequence.
AC022408
AC022408
AC022408.6 GI:15217198
HTG.
SOURCE
Homo sapiens (human)
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniota; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Hominae; Homo.
REFERENCE
1 (bases 1 to 150266)
Rowen,L., Madan,A., Qin,S., Abbasi,N., Baradarani,L., Birditt,B., Bloom,S.,
Burke,J., Dors,M., Fleetwood,P., Kaur,A., Madan,A., Nesbitt,R.,
Pate,D., and Hood,L.
Sequencing of human chromosome 15 D15S146-D15S117 region
Unpublished
2 (bases 1 to 150266)
Rowen,L., Madan,A., Qin,S., Abbasi,N., Baradarani,L., Birditt,B.,
Bloom,S., Dors,M., Dichtoff,R., Fleetwood,P., Harrison,G.,
James,R., Kaur,A., Madan,A., Owen,M.P., Ratcliffe,A., Shaffer,T.,
and Hood,L.
Direct Submission
Submitted (03-FEB-2000) Multimegabase Sequencing Center, University
of Washington, PO BOX 357730, Seattle, WA 98195, USA
3 (bases 1 to 150266)
Rowen,L., Madan,A., Qin,S., Baradarani,L., Birditt,B., Bloom,S.,
Burke,J., Dors,M., Fleetwood,P., Kaur,A., Madan,A., Nesbitt,R.,
Pate,D., and Hood,L.
Direct Submission
Submitted (21-AUG-2001) Multimegabase Sequencing Center, Institute
for Systems Biology, 4225 Roosevelt Way NE, Suite 200, Seattle, WA
98105, USA
On Aug 21, 2001 this sequence version replaced gi:1125361.
----- Genome Center
Center: Multimegabase Sequencing Center
Center code: UWMSC
Web site: http://chroma.mbc.washington.edu/msg_www
Contact: leetown@systemsbiology.org
----- Summary Statistics
Sequencing vector: pUC18; 108752
Chemistry: Dye-terminator Big Dye; 90% of reads
Assembly program: Phrap; version 0.990399
Note: data from AC021753 [Drafting center UWMSC] and AC012652
[Drafting center UWMSC] and 58017 [Drafting center WUGSC] were
added for finishing.
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overlapping BACs were combined and the consensus sequence
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/note="Low quality data."
100628..150266
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ORIGIN

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Chemistry: Dye-primer Big Dye; 10% of reads
Assembly program: Phrap; version 0.99039
Note: data from AC022408 (Drafting center UMSC) and AC020661
were added for finishing.

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ORIGIN

Query Match 18.4%; Score 459.4; DB 8; Length 167996;
Best Local Similarity 62.0%; Pred. No. 5.8e-83;
Matches 888; Conservative 1; Mismatches 487; Indels 57; Gaps 8;

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119935 A--TTTTCTGTAGAGACAGGAGTCTTCAAGTTGGCCAGGCTTGTCTTAACTCTG 119992
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120231 CCTCTGAGTGTGAGATTTAACAAGGCTGACCAACCATCTGCTAATTTTGTATTT 120290
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120291 TTGTTAGATGAGATTTAACAATGTTGGCAGGCTGTTTCAAACTCTGACCTCAAGT 120350
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1797 GAGAACTTGAAGACAGGCTGCTGATTAATTAATTAATTAATTAATTAATTAATTAATTA 1856
120901 CAGGGGCTGAGACAGGCTGACCAACCAATTAATTAATTAATTAATTAATTAATTAATTA 120960
1857 AATTAAGTGGGTGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 1916
120961 AATTAAGTGGGTGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 121020
1917 AGAATGCTTGAACCTGAGGAGGACAGAGGTTGACAGTGAAGTCCAGATCCACATGCACTC 1976
121021 AGAATGCTTGAACCTGAGGAGGACAGAGGTTGACAGTGAAGTCCAGATCCACATGCACTC 121080
1977 CAGCTGAGGACACAGGACGATCTATCTCAAAAATTAATTAATTAATTAATTAATTA 2029
121081 CAGCTGAGGACACAGGACGATCTATCTCAAAAATTAATTAATTAATTAATTAATTA 121133

RESULT 11
AC003689 137693 bp DNA linear PRI 21-NOV-1998
LOCUS Homo sapiens Chromosome 11q12.2 PAC clone pJ1081b4 containing
DEFINITION human mRNA for T-cell glycoprotein CD6, complete sequence.
ACCESSION AC003689
VERSION AC003689.1 GI:3900834
KEYWORDS HTG.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Hominidae; Homo
1 (bases 1 to 137693)
REFERENCE
Evans G.A., Athanasiou M., Aguayo P., Armstrong D., Basit M.,
Buetner J., Bumeister R., Card P., desallbois F., Dunn J.,
English C., Ehrhard S., Garner H.R., Gee V., Gordon M., Gotway G.,
Grant O., Hahner L., Joslin J., Lewis B., Loo H., Loo K.N.,
Major T., McFarland J., Newton J., Osborne-Lawrence S.,
Schegeman J., Schultz R.A., Stimson S., Syed M. and Ward T.
HTGS Submission
Unpublished

REFERENCE 2 (bases 1 to 137693)
AUTHORS Evans, G.A., Athanasiou, M., Basit, M., Bradbury, P., Brignac, S., Bunesler, R., Davis, C., English, C., Franklin, T.L., Garner, H.R., Gee, V., Gordon, M., Gotway, G., Grant, O., Hahner, L., Harris, J., Hinson, S., Narayanaswamy, U., Newton, J., O'Brien, K., Patel, P., Schageman, J., Schilling, P., Schultz, R., Syed, M., Valenzuela, D., Ward, T. and Wilson, R.
TITLE Direct Submission
JOURNAL Submitted (17-DEC-1997) Genome Science & Technology Center, University of Texas Southwestern Medical Center, 5323 Harry Hines Blvd, Dallas, TX 75235-8591, USA
REFERENCE 3 (bases 1 to 137693)
AUTHORS Evans, G.A., Athanasiou, M., Aguayo, P., Armstrong, D., Basit, M., Bunesler, R., Butler, C., Card, P., Desaliboat, F., Dunn, J., English, C., Ethridge, S., Garner, H.R., Gee, V., Gordon, M., Gotway, G., Grant, O., Hahner, L., Joslin, J., Lewis, E., Loo, H., Loo, K.N., Major, T., McFarland, J., Newton, J., Osborne-Lawrence, S., Schageman, J., Schultz, R.A., Stimson, S., Waller, K. and Ward, T.
TITLE Direct Submission
JOURNAL Submitted (21-NOV-1998) Genome Science & Technology Center, University of Texas Southwestern Medical Center, 5323 Harry Hines Blvd, Dallas, TX 75235-8591, USA
COMMENT On Nov 21, 1998 this sequence version replaced gi:2695567. Further information regarding the map of this region or annotation of pDJ1081b4 can be found at <http://gsstec.swmed.edu/chromosol.htm>
IMPORTANT: This submission contains the entire insert of clone pDJ1081b4. pDJ1081b4 comes from the RPCR-3 PAC library constructed at the Roswell Park Cancer Institute by the Pletier de Jong group. This clone has been finished according to strict quality criteria and attempts have been made to resolve all base calling problems such as compressions and repetitive elements. The expected phred/Phrap calculated errors/10kb is 0.98. In addition, attempts have been made to assure over 99% of consensus base calls consist of either double-stranded coverage or 2 types of labeling chemistry on one strand.
CHROMOSOMAL LOCUS: This PAC clone comes from the Chromosome 11p12.2 Bear's disease region mapped between STS D11S461 and EST AHNAK. This region spans over 1.5 Mbp.
MARKER CONFIRMATION: ESTs/STS sequence confirmed, CD6, D11S461
MAPPED CLONE OVERLAP: PACs pDJ6066 and pDJ59421.
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repeat_region
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/rpt_family="Alu"
repeat_region


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VERSION      AP003721.3  GI:31790667
KEYWORDS     HTG.
SOURCE       Homo sapiens (human)
ORGANISM     Homo sapiens
              Eukaryota; Metazoa; Chordata; Craniata; Vertebrate; Euteleostomi;
              Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
              Homnidae; Homo.
REFERENCE    1
AUTHORS      Hattori, M., Ishii, K., Toyoda, A., Taylor, T.D., Hong-Seog, P.,
              Fujiyama, A., Yada, T., Totoki, Y., Watanabe, H. and Sakaki, Y.
TITLE        Homo sapiens genomic DNA
JOURNAL      Published Only in Database (2001)
REFERENCE    2 (bases 1 to 201460)
AUTHORS      Hattori, M., Ishii, K., Toyoda, A., Taylor, T.D., Hong-Seog, P.,
              Fujiyama, A., Yada, T., Totoki, Y., Watanabe, H. and Sakaki, Y.
TITLE        Direct Submission
JOURNAL      Submitted (04-JUN-2001) Masahira Hattori, The Institute of Physical
              and Chemical Research (RIKEN), Genomic Sciences Center (GSC);
              1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
              (E-mail: hattori@gsc.riken.go.jp, URL: http://hgp.gsc.riken.go.jp/,
              Tel:81-45-503-9111, Fax:81-45-503-9170)
              On Jun 16, 2003 this sequence version replaced gi:28189463.
COMMENT      Location/Qualifiers
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Best Local Similarity 59.7%; Pred. No.2.5e-82;
Matches 877; Conservative 1; Mismatches 574; Indels 17; Gaps 6;
QY      604 TTTTGTGTTGTTGTTTGAAGACAGGGCTTGTCTC-TGTACCCAGGAGTGAAGACAG 662
DB      144460 TTTCTTTTGTGTTTGTGTTTGAAGACAGGTTTCTCTGTTGGCCAGGCTGGAGTGAA 144519
QY      663 TGTGCAACCAVAGGTCACTGCAAGCTCAACCTCTGAGCTCAAGAGATCTGTGACCTG 722
DB      144520 TGGCAACATCTGGGTTCACCAACCTCCGCTCCGGGGTTCAAGGGTTTCTCCGCTC 144579
QY      723 AGCTTCCCAATAGCTGGGAGTCAAGAGGTGACACACAGCGCTGGCTAATTAATAAAT 782
DB      144580 AGCTCCCGAGTACGTGAGATTAACAGGATGCGCACACGCCAGCTGATTTTGTGA--- 144636
QY      783 TTTTGTGAGAGACTGGGCTTTACTAGTGTGGCAGGCTTGTCTTAATCTCTGGCTTCA 842
DB      144637 TTATTAGTAGAGAGGGGGTTTCTCATGTTGTGTCAAGCTTGAGTCTTGAGCTCTCA 144696
QY      843 AGCAATCTCTTACCTTGGATCCCAAGTGTGGGATTAACAGGGGTGAGCCACCATGTG 902
DB      144697 GGTATCCACCCACCTCGGCTCCCAAGTGTGGGATTTAGGTGTAGGCGCATCAGGCC 144756
QY      903 CGGCTACTTATTTCTTTCATTTCACTTCTTCAATAGATGTAAATTCACAGAACAGGG 962
DB      144757 CGGGCTCAACATTTCTTAATCTTAATGTTTTCACGCTTGAAGTCTCAAGTATATCA 144816
QY      963 ATTACTGCTTATTTCTCTCTTCTTTTGAAGACAGAGTCTCACTTCAACCTCAACC 1022
DB      144817 CTTTGATTTCTTTTGTGAGACAGTCTTGCTCTGTGTCAACAGGCTGAGAGTCAATGCG 144876
QY      1023 TCCGTTAGCTCACTGCAACCTGTGCTCCGGGTTCAAGGATTTCTCTGCTTAAGCTT 1082
DB      144877 CAATCTCGGCTCACTGCAAACTGTGCTCCGAGGTTCACGGATTTCTCTCCTCAGGCT 144936
QY      1083 CCGTAGTAGCTGGAATTAACAAGCGTGAACCAACATGCTTGGCTAATTTT--GTAATTTT 1140
DB      144937 CCAAGTAGCTGGGATTAACAAGTGGGCCCAACAGTCTGCTACTTTTGTATATTTT 144996
QY      1141 TAGCAGAGATGGGGTTTATCCATGTTGCCAGGCTGTGTCTCAAACTCTGAACCTCAAGTG 1200

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DB      145057 ATTCGCTGCTCTGAGTCTCCCAAGTGTGGAATTTATAGGGGTAGTACGACCGGCCAT 145116
QY      1261 CGATTACTGCTATTTTCTTAATGTGCTATATCCCAAGTCTAGAGAGTGTCTGACATAT 1320
DB      145117 GCCCTTGATTTCTTTGAACAAATGTAATTAATTAATTAATTAATTAATTAATTAATTAAT 145175
QY      1321 AGTAGTGTCTCAATTAATTAATTAATTAATTAATTAATTAATTAATTAATTAATTAATTA 1380
DB      145176 CCTGTGATCAATTAATTAATTAATTAATTAATTAATTAATTAATTAATTAATTAATTA 145235
QY      1381 TTTTAAACATATCTTGACAACTTTGACAAATTAATTAATTAATTAATTAATTAATTAATTA 1440
DB      145236 ATTCACACATTAACCAAAATTAATTAATTAATTAATTAATTAATTAATTAATTAATTA 145295
QY      1441 C-----TTATCACTTGTATGACTTTTTCATATGCTCAAACTTAATTTGTTAC 1491
DB      145296 ATATAAGATGATTTAGCAGAGAAACTAATGTTATTTGAAAGATATCTGAATGGGTG 145355
QY      1492 TGTTTTTCATTTGTTACTATTTTATGATCACTGAATTAATTAATTAATTAATTAATTAATTA 1551
DB      145356 TTTAGTTTTTCCCTCTCACTTATGTAACATACCAGAGATGATATTAATTTGTTTCTAG 145415
QY      1552 CTCTGCTCACTTTAGAGGCCAAATTTACAAATCTGATGAAGTAACTGTAACCTCTCC 1611
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QY      1791 TGAAGTAGAAGTGTGAGACCAAGCTGTGTAATTAATTAATTAATTAATTAATTAATTAATTA 1850
DB      145656 TGAAGTAGAAGTGTGAGACCAAGCTGTGTAATTAATTAATTAATTAATTAATTAATTAAT 145715
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QY      2031 AGATTCGAGAGAAACAAACTAATAAGA 2059
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RESULT 13
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LOCUS     Papio anubis clone RP41-1C24, WORKING DRAFT SEQUENCE, 4 ordered
DEFINITION
pieces.
AC116933
AC116933.2 GI:27356688
VERSION  HTG; HTGS PHASE2; HTGS DRAFT.
KEYWORDS Papio anubis (olive baboon)
SOURCE

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Oy 1130 TTTGATTTTGTAGCAGAGATGGGGTTTTTACATGTTGCCAGGCTGGTCTCAACTCT 1189
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Oy 1885 GTAATCCAGCTACTCGGAGGCTGAGGCAAGAGATTTGTTGAACCTTGGAGGAGAGG 1944
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RESULT 14
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 ACCESSION AC005052
 VERSION AC005052.2 GI:10122134
 KEYWORDS HTG.

SOURCE
 ORGANISM Homo sapiens (human)
 Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
 Homnidae; Homo.
 REFERENCE
 AUTHORS Sulston, J.E. and Waterston, R.
 TITLE Toward a complete human genome sequence
 JOURNAL Genome Res. 8 (11), 1097-1108 (1998)
 PUBMED 9847074
 REFERENCE
 AUTHORS Tsin-Mollam, A., Graves, T. and Colman, M.
 TITLE The sequence of Homo sapiens BAC clone CTB-38K21
 JOURNAL Unpublished
 REFERENCE
 AUTHORS Waterston, R.H.
 TITLE Direct Submission
 JOURNAL Submitted (12-JUN-1998) Genome Sequencing Center, Washington
 University School of Medicine, 4444 Forest Park Parkway, St. Louis,
 MO 63108, USA
 REFERENCE
 AUTHORS Waterston, R.H.
 TITLE Direct Submission
 JOURNAL Submitted (14-SEP-2000) Genome Sequencing Center, Washington
 University School of Medicine, 4444 Forest Park Parkway, St. Louis,
 MO 63108, USA
 REFERENCE
 AUTHORS Waterston, R.
 TITLE Direct Submission
 JOURNAL Submitted (08-NOV-2000) Department of Genetics, Washington
 University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
 On Sep 14, 2000 this sequence version replaced gi:3212939.
 COMMENT
 Genome Center
 Center: Washington University Genome Sequencing Center
 Center code: WUGSC
 Web site: http://genome.wustl.edu/sec
 Contact: saplens@watson.wustl.edu
 ----- Summary Statistics
 Center project name: H_RG038K21

NOTICE: This sequence may not represent the entire insert of this
 clone. It may be shorter because we only sequence overlapping
 clone sections once, or longer because we provide a small overlap
 between neighboring data submissions.

This sequence was finished as follows unless otherwise noted:
 all regions were double stranded, sequenced with an alternate
 chemistry, or covered by high quality data (i.e., phred quality >=
 30); an attempt was made to resolve all sequencing problems, such
 as compressions and repeats; all regions were covered by sequence
 from more than one subclone; and the assembly was confirmed by
 restriction digest.

MAPPING INFORMATION:
 This sequence was generated from part of bacterial clone contigs of
 human chromosome X, constructed by the chromosome X mapping group
 at the Sanger Centre, Wellcome Trust Genome Campus, Hinxton, UK.
 Further information can be found at
 http://www.sanger.ac.uk/HGP/ChrX/

SOURCE INFORMATION:
 Clone CTB-38K21 is from the first release of the human BAC library
 CTB-978SK-B. The library contains cloned DNA from the male
 fibroblast cell line 978SK. See: Shizuya et al., Proc. Natl.
 Acad. Sci. USA 89:8794-7 (1992); U-J. Kim et al., Genomics 34:213-8
 (1996). This clone is available from Research Genetics, Inc.
 (http://www.resgen.com).
 VECTOR: pBel0BAC11
 Selection: chloramphenicol

NEIGHBORING SEQUENCE INFORMATION:
 The clone sequenced to the right is RP3-327A19, 200 base pair

overlap. Actual start of this clone is at base position 1 of
CTB-38K21, actual end is at base position 9416 of RP3-327A19.
Location/Qualifiers

FEATURES

SOURCE

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repeat_region
12405..12520
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repeat_region
12712..13008
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repeat_region
13009..13315
/rpc_family="Alu"
repeat_region
13603..13850
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13851..14147
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repeat_region
14148..14222
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repeat_region 14527..14713
/rpc_family="ERV4"
repeat_region 14815..15244
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repeat_region 15245..15566
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repeat_region 16447..16580
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repeat_region 17255..17390
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repeat_region 21559..21803
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21835..21951
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Query Match 18.0% Score 449.2; DB 8; Length 134210;
Best Local Similarity 59.5%; Pred No.6.9e-81;
Matches 896; Conservative 1; Mismatches 554; Indels 55; Gaps 6;

QY 555 TGAGCTATTTGGCCAAATCACACAGCTTGAAGTGTGACAGTTGGCTTTTGTCT 614
DB 53793 TGTGGATTTGCTTATTATTATTATTATTTTGTGTTAGGGGCTGATTCCTTTT 53734
QY 615 TGTGTTTAGACACAGGCTTGTCTGTGCACCCAGGAGTATGACACAGTGTGACAT 674
DB 53733 TCTTTTGTGACACAGGCTTCTCTGTGCACCCAGGCTGTGAGTGTGACAC-ATCAT 53675
QY 675 AGGTACGTGACAGCTCAACCTCCTAGCTCAAGGATGTGACCTCAGCTCCCAAGT 734
DB 53674 GGCTCAGTCAAGCTTCAACCTTCTGGCTCAAGCAATCTCAAAATCTCAGCTCCGAGT 53615
QY 735 AGCTGGACTACAGAGGTGACACACACAGCTGGCTAATTTAAAAATTTTGTAGAG 794
DB 53614 AGCTGGGTCTAATAGGCTTAGGTGACACAGGCCGGCTAATTTTAAATGTTTGTAGAGA 53555
QY 795 ACTGGGTCTTACAGTGTGACAGGCTTGTCTTAACTCTGGCTTCAAGCAATCTCTCT 854
DB 53554 CAAGGGTCTCAGTATGTACACAGGAGGTGTCTTAAATCTCGGTTCAAGGAGATCTCTCT 53495
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OY	855	ACCTGGGATATCCAAAGGCTGGGATTAACAGGGGTGAGCCACATGCGGGCTACTATT	914
Db	53494	GCCTTGGCTCCAAAGCTGAGATTACAGGATGAGCCACTGCACCGGCTGATTCC	53435
OY	915	TCCTTACATTCGATCTTTCCATTAAGATGTAAATCCAGAAACAGGATTACTGCTAT	974
Db	53434	TGATTTCTTTTCTTTT-----TTCTTCT	53409
OY	975	TTTCTCTCTTTCTTTTGTAGACAGATCTCACTTCACTCCACCTCCGTTCA----	1030
Db	53408	TTTCTTTTTTTTTTTTTGTAGACGGAATCTCGCTCTGTCCGACGAGCTGAGAGCTGTG	53349
OY	1031	-----GCTCACTGCAACCTCTGCTCCCGGGTTCAAGTATCTCTGCTCAAGC	1080
Db	53348	CAGACTTGTGCTCACTGCAAGGTCCACTCTCTGGGTTACGCGCATTTCTCTGCTCACG	53289
OY	1081	CTCTGAGTAGCTGGAAATTACAGCGTGCACACCATGCTGGCTAATTTTTGTATTTT	1140
Db	53288	CTCGGAGTAGCTGGGACTACAGGTGCCCCCACCACACCCACACTAAATTTTTGTATTTT	53229
OY	1141	TAGCAGAGATGGGGTTTTTCAATGTTGCCAGGCTGGTCTCAACCTCGACTCAAGT	1200
Db	53228	TAGTAGAGACGGGGTTTTCAACGGTGTAGCAGAGATGTCTCGATCTCTGACCTC--GTG	53171
OY	1201	ATCTGCTGCTCAGTCTCCCAAAGCTGGAATTATAGCGGTGATCACTGTGCTGCG	1260
Db	53170	ATCGGCTGCTCGGCTCCCAAAGTCTGGGATTTACAGCATGAGCCACCGGCTGCG	53111
OY	1261	CGATTACTGTCTATTTTCTTATTTGCTATATCCCGAATCTTAGAGCAGTCTGCATAT	1320
Db	53110	CTTTTCTCTTTTATTTTTTTTTGTAGACAGGGCTTC--CTCTGTCCACCAGCGTATATG	53052
OY	1321	AGTAGTGCTCAATAAATAATGATGATGCACAGCTAGATATAACTTTCTTTTCTT	1380
Db	53051	ACTGGCACTATCTTGCTCACTGCACCTCACTTCCAGGTTCAAGCAATTATCATGCG	52992
OY	1381	TTTTTAAACAATCTTGCACAACTTTCGAAATTAATCAATCTTGCAATTCGCTTTTCA	1440
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OY	1441	CTTATCACTTGTATATGACTTTTTCATATGCTCAAAAC---CTTATATGTAAGTTTT	1497
Db	52931	TTTTTATGAAGAGAGGTTTCCCACTGTGGCGAAGATGTCCTCATCTTGAACCTG	52872
OY	1498	TTCAATGTACTATTTTATGCTAGATATATATGCTTATTTGCTTATATACATCCCTG	1557
Db	52871	TGATCTGCGCGCTCGGCTCCCAAAGTGTGAGATTACAGGGTAGGCACTGTGCCG	52812
OY	1558	CTCAGCTTTAGAGGCCAAATTTACAAATCTGATGAAGCTATGAACCTCTGCCAGAG	1617
Db	52811	GCCGATPAACAGCTTTTATATGTGCACATCTGCTCAAGTTATATGTATATATAT	52752
OY	1618	AAATACACACACACACACTCACACAGTTTTTTTTTATATGTTTGAACATAAGACA	1677
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OY	1678	AGAAACCTGCATTAGAGATGTTTGTCAATATTAATTAATAATATCACTAGTTGGGCAG	1737
Db	52691	ACCTGCTGATTAAGAATCACTACTGAGACTAGAAAGAAAGATGTTTAGGCCAGGAGTGG	52632
OY	1738	TGACTCAAGCTGTAAACACAGTACTTTTGAAGTTCACAGGTGGGTGATCACTGAGGTG	1797
Db	52631	TAGCTCACTGTATATCCAGCACTTTGGAGGCCGAGGTGGAAGATTAACCTGAGATC	52572
OY	1798	AGAAATTGAGACCAAGCTGTGATATATGTTGAACCTATCTTATCTAATAATACAAA	1857
Db	52571	AGGAGTTTGAAGACCACTGACCAATGGCAAAACCCATCTCTATAAATAATCAAAA	52512
OY	1858	ATTAGCTGGGTATAGTATGATGATCCCTGTATGTCCAGCTACTTGGGAGGCTGAGGCAGA	1917
Db	52511	ATTATGAGGGATGTGTATCATATGCTGTGATCTCAGCTACTTGGGAGGCTGAGGCAGA	52452

QY	1918	GAATTGCTTGAACCTGGAGGCGAGAGTTGCAGTGAAGCCGAGATCCACCACTGCATCC	1977
QY <td>1918 <td>GAATTGCTTGAACCTGGAGGCGAGAGTTGCAGTGAAGCCGAGATCCACCACTGCATCC</td> <td>1977</td> </td>	1918 <td>GAATTGCTTGAACCTGGAGGCGAGAGTTGCAGTGAAGCCGAGATCCACCACTGCATCC</td> <td>1977</td>	GAATTGCTTGAACCTGGAGGCGAGAGTTGCAGTGAAGCCGAGATCCACCACTGCATCC	1977
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QY	1978	AGCCTGGGCGACACAGCGAGACTCTATCTCAAAAAATAATTAATAATAAGATCG	2037
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QY	2038	GAGAGA 2043	
Db	52331	AAAAAA 52326	

RESULT	LOCUS	DEFINITION	ACCESSION	VERSION	KEYWORDS	SOURCE
15	AC005484.5	131943 bp	DNA	linear	PRI 30-SEP-2000	
	AC005484	Homo sapiens	PAC clone RPS-84708	from 14q24.3,	complete sequence.	
	AC005484	AC005484.2	GI:5091654			
	HTG.	Homo sapiens (human)				

REFERENCE 1 (bases 1 to 131943)

AUTHORS Sulston, J. E. and Waterston, R.
TITLE Toward a complete human genome sequence
JOURNAL *Nature* 343 (1990) 319-321

JOURNAL
PUBMED
Genome Res. 8 (11), 1097-1108 (1998)
9847074

REFERENCE	2 (bases 1 to 131943)
AUTHORS	Cloud, J., Wohlmann, P. and Holmes, A.

TITLE The sequence of homo sapiens PAC clone KFS-84/08
 JOURNAL Unpublished
 ORIGINATOR (b) (5) (11043)
 ORIGINATOR 3

REFERENCE	3 (pages 1 to 131943)
AUTHORS	Waterston, R.H.
TITLE	Direct Substitution

Submitted (14-AUG-1998) Genome Sequencing Center, Washington University School of Medicine 660 South Euclid Ave. St. Louis, MO 63110

UNIVERSITY SCHOOL OF MEDICINE, 5555 FOR
MO 63108, USA
(bases 1 to 131943)
REFERENCE

REFERENCE	4 (pages 1 to 15,745)
AUTHORS	Waterston, R.H.
TITLE	Direct Submission

JOURNAL
Submitted (17-JUN-1999) Genome Sequencing Center, Washington
University School of Medicine 4444 Forest Park Parkway St Louis

CARLEIGH, DONALD OF HOUSTON, 1111 101000 MAIN BLVD., SU. 2000
 MO 63108, USA
 5 (bases 1 to 131943)
 PREFERENCE

AUTHORS Waterston, R.
TITLE Direct Submission

Submitted (30-SEP-2000) Department of Genetics, Washington University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA

COMMENT On Jun 17, 1999 this sequence version r
----- Genome Center

Center: Washington University Genome Sequencing Center
Center code: WUGSC

Web site: <http://genome.wustl.edu/gsc>
Contact: sapiens@watson.wustl.edu

----- Summary Statistics
Center project name: H DJ0847008

.....

NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping.

clone sections once, or longer because we provide a small overlap between neighboring data submissions.

This sequence was finished as follows unless otherwise noted:

all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phased quality >=

as compressions and repeats; all regions were covered by sequence fragments that are subcloned and the assembly was confirmed by

from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:
This clone from chromosome 14 was provided by Dr. Pieter de Jong, Roswell Park Cancer Institute, Human Genetics Department, Elm and Carlton Streets, Buffalo NY 14263-0001 USA.

SOURCE INFORMATION:
This clone was derived from human PAC library RPCI-5, prepared by Pieter de Jong and coworkers at the Roswell Park Cancer Institute (http://pacpac.med.buffalo.edu) using the method described by Ioannou et al., Nature Genetics 6:84-9 (1994). The library is from one male donor.

The clone may be obtained either from Genome Systems, Inc. (http://www.genomesystems.com) or Research Genetics, Inc. (http://www.resgen.com) or from Pieter de Jong.

VECTOR: pCYPAC2

NEIGHBORING SEQUENCE INFORMATION:

The clone sequenced to the left is RP5-89265; the clone sequenced to the right is RP4-59267, 200 bp overlap. Actual start of this clone is at base position 1 of RP5-84708; actual end is at base position 131747 of RP5-84708.

Location/Qualifiers

1..131943

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/organism="Homo sapiens"

/mol_type="genomic DNA"

/db_xref="taxon:9606"

/chromosome="14"

/map="14q24.3"

/clone="RP5-84708"

/clone_1lb="RPCI-5"

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1762..2225

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3651..3916

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3917..3943

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10890..11242

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11567..11996

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12006..12113

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12123..12146

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12147..12430

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12445..12535

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12527..13097

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13217..13339

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14138..14158

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16569..16963

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/rpt_family="Alu"

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Query Match 17.9%; Score 447.8; DB 8; Length 131943;

Best Local Similarity 61.3%; Pred. No. 1.3e-80; Indels 52; Gaps 10;

Matches 913; Conservative 1; Mismatches 523;

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GenCore version 5.1.7
Copyright (c) 1993 - 2006 Bioceleration Ltd.

OM nucleic - nucleic search, using sw model

Run on: February 17, 2006, 10:20:55 ; Search time 9872 Seconds
(without alignments)
1185.170 Million cell updates/sec

Title: US-10-607-806-1-C7256_COPY_7000_9500
Perfect score: 249
Sequence: 1 gtcgtgcaactgctgcacg.....ttcgagaccagcctgacaa 2501

Scoring table: IDENTITY_NUC
Gapop 10.0, Gapext 1.0

Searched: 41078325 seqs, 23393541228 residues

Total number of hits satisfying chosen parameters: 82156650

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%

Listing first 45 summaries

Database :

EST.*
1: gb_est1.*
2: gb_est2.*
3: gb_est3.*
4: gb_est4.*
5: gb_est5.*
6: gb_est6.*
7: gb_est7.*
8: gb_est8.*
9: gb_est9.*
10: gb_est10.*
11: gb_est11.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	505.4	20.2	507	BU785040	in43909.x
2	344.2	13.8	4087	BC024593	BC024593 Homo sapi
3	341	13.6	3990	AL713681	AL713681 Homo sapi
4	337.2	13.5	15970	AO839852	AO839852 260L13-C5
5	315.8	12.6	897	CD557847	CD557847 AGENCOURT
6	311	12.4	5797	CR858326	CR858326 Pongo pyg
7	307.2	12.3	5797	CR749233	CR749233 Homo sapi
8	301.2	12.1	1605	CR860521	CR860521 Pongo pyg
9	301.2	12.1	1863	CR599842	CR599842 full-leng
10	300.6	12.0	946	BO958903	BO958903 AGENCOURT
11	298	11.9	918	BO706343	BO706343 AGENCOURT
12	293.6	11.7	736	CK780936	CK780936 HES3_2_B
13	290.6	11.6	3092	CR613629	CR613629 full-leng
14	288.2	11.5	855	BO681302	BO681302 AGENCOURT
15	287.6	11.5	769	BO710315	BO710315 AGENCOURT
16	285.2	11.4	4088	BS483111	BS483111 Homo sapi
17	283	11.3	3165	CR859576	CR859576 Pongo pyg
18	281.6	11.3	800	AU120942	AU120942 Pongo pyg
19	280.4	11.2	672	CA431692	CA431692 UI-H-DFO-
20	280	11.2	3474	AL110229	AL110229 Homo sapi
21	278.8	11.2	658	AO393450	AO393450 CITBI-E1-
22					

23	277.6	11.1	877	9	AQ739838	AQ739838 HS_5505_A
24	277.2	11.1	1875	4	BC009270	BC009270 Homo sapi
25	276.6	11.1	3990	4	BSM803026	BSM803026 Homo sapi
26	276.4	11.1	4828	4	BSM802759	BSM802759 Homo sapi
27	276.2	11.1	881	5	BU521286	BU521286 AGENCOURT
28	274.8	11.0	3552	4	CR860263	CR860263 Pongo pyg
29	273.8	11.0	736	6	CA427039	CA427039 UI-H-DFO-
30	273.4	10.9	629	5	BS509360	BS509360 DKFZP686J
31	273.4	10.9	2230	4	CR859082	CR859082 Pongo pyg
32	272.4	10.9	652	6	CA427045	CA427045 UI-H-DFO-
33	272.4	10.9	666	6	CA431783	CA431783 UI-H-DFO-
34	272.4	10.9	815	8	CK785622	CK785622 AGENCOURT
35	272.2	10.9	922	5	BU501973	BU501973 AGENCOURT
36	272	10.9	617	3	BI861844	BI861844 603388872
37	271.8	10.9	644	7	CN480313	CN480313 UI-H-EUO-
38	271.6	10.9	1042	5	EX377759	EX377759 BX377759
39	271.6	10.9	2097	4	CR614786	CR614786 full-leng
40	271.4	10.9	603	1	AL707313	AL707313 DKFZP686P
41	271.2	10.9	5325	4	BSM80409	BSM80409 Homo sapi
42	271.2	10.9	5785	4	BSM802309	BSM802309 Homo sapi
43	270.6	10.8	2330	4	CR615928	CR615928 full-leng
44	270.4	10.8	618	9	BZ609884	BZ609884 WACW767F
45	270	10.8	665	5	BU633001	BU633001 UI-H-DFO-

ALIGNMENTS

RESULT 1
LOCUS BU785040 507 bp mRNA EST 11-OCT-2002
DEFINITION in43909.xl HR85 16let Homo sapiens cDNA clone IMAGE:6125008 3',
mRNA sequence.
ACCESSION BU785040
VERSION BU785040.1 GI:23830576
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Bukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Homidae; Homo.
1 (bases 1 to 507)
Melton,D., Brown,D., Kenty,G., Permutt,A., Lee,C., Kaestner,K.,
Lemishka,I., Scaerce,M., Brestelli,J., Gradwohl,G., Clifton,S.,
Hillier,L., Marra,M., Pape,D., Wylie,T., Martin,J., Birstein,A.,
Schmitt,A., Theising,B., Ritzer,B., Komko,I., Bennett,U.,
Cardenas,M., Gibbons,M., McCann,R., Cole,R., Tsagarisvilli,R.,
Williams,T., Jackson,Y. and Bowers,Y.
Endocrine Pancreas Consortium
Unpublished (2000)
Contact: Douglas Melton, Klaus H. Kaestner, & Hiroshi Inoue
Endocrine Pancreas Consortium
Harvard University, Howard Hughes Medical Institute
Dept of Molecular and Cellular Biology, 7 Divinity Ave, Cambridge,
MA 02138
Tel: 617-495-1812
Fax: 617-495-8557
Email: dmelton@biochem.harvard.edu
Library was constructed by Dr. Hiroshi Inoue DNA sequencing by:
Washington University Genome Sequencing Center For information on
obtaining a clone please contact: Dr. Hiroshi Inoue
(hinoue@im.wustl.edu)
Seq primer: -40UP from Gibco
High quality sequence stop: 443.
Location/Qualifiers
1. 507
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:6125008"
/tissue_type="Purified pancreatic islet"
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/clone_1ib="HR85 16let"

FEATURES

source

/note="Organ: Pancreas; Vector: pBluescript SK(-); Site_1: NotI; Site_2: XhoI; cDNA made by oligo-dT priming. Size selected on agarose gel. Average insert size ~1kb. 5' XhoI site was destroyed after directional cloning. Amplified once. Contact Information: Hiroshi Inoue, MD, Metabolism Div. (Alan Permut Lab), Washington University School of Medicine, Box 8127, 660 South Euclid Ave., St. Louis, MO 63110, E-mail: hinooue@ingate.wustl.edu, Tel: 314-362-1916, Fax: 314-747-2692."

ORIGIN

Query Match 20.2%; Score 505.4; DB 5; Length 507;
Best Local Similarity 99.8%; Pred. No. 2.3e-35;
Matches 506; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1506 TACTATTATTAGTCACTGTAATAATATGCTTAATTTGCTTAATACATCCCTCCCTCCACTT 1565
DB 507 TACTATTATTAGTCACTGTAATAATATGCTTAATTTGCTTAATACATCCCTCCCTCCACTT 448
QY 1566 TAGAAGGCCAATTATACAAATCTGATGTAAGCTATGAAACCTCTCCCGCAGAAATACAC 1625
DB 447 TAGAAGGCCAATTATACAAATCTGATGTAAGCTATGAAACCTCTCCCGCAGAAATACAC 388
QY 1626 ACACACACACACTCACAACAGTTTTTTTTTAAATGTTTSCAACTAAGACAGAAACCT 1685
DB 387 ACACACACACACTCACAACAGTTTTTTTTTAAATGTTTSCAACTAAGACAGAAACCT 328
QY 1686 GCATTAGAGATGTTTGTTCATATTAATAAATACTAGTTGGGACAGTACCTCA 1745
DB 327 GCATTAGAGATGTTTGTTCATATTAATAAATACTAGTTGGGACAGTACCTCA 268
QY 1746 GCCTGTAAACACAGTACTTGTGAAGTCCAGGTGGGTGATCACTTGAGTGAAGAATTC 1805
DB 267 GCCTGTAAACACAGTACTTGTGAAGTCCAGGTGGGTGATCACTTGAGTGAAGAATTC 208
QY 1806 GAGACACAGCTGTGCAATATGTTGAACCCCTATCTCTAATAAATACTAATAATTAAGTGTG 1865
DB 207 GAGACACAGCTGTGCAATATGTTGAACCCCTATCTCTAATAAATACTAATAATTAAGTGTG 148
QY 1866 GGTGTAGATGATGATGCTGCTGATGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 1925
DB 147 GGTGTAGATGATGATGCTGCTGATGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 88
QY 1926 TGAACCTGAGAGGAGAGGAGTGTGAGTGAAGGAGATCCCACTGCACTCCAGCTGGG 1985
DB 87 TGAACCTGAGAGGAGAGGAGTGTGAGTGAAGGAGATCCCACTGCACTCCAGCTGGG 28
QY 1986 CGACACAGCAGAGACTCTATCTCAAAA 2012
DB 27 CGACACAGCAGAGACTCTATCTCAAAA 1

RESULT 2
BC024593 4087 bp mRNA linear HTC 29-JUN-2004
LOCUS Homo sapiens cDNA clone IMAGE:3914314, with apparent retained
DEFINITION intron.
ACCESSION BC024593
VERSION BC024593.1 GI:22137609
KEYWORDS HTC.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Homnidae; Homo.

REFERENCE
AUTHORS
1 (bases 1 to 4087)
Strausberg, R.L., Feingold, E.A., Grouse, L.H., Derge, J.G.,
Klausner, R.D., Collins, F.S., Wagner, L., Shenmen, G.D.,
Altschul, S.F., Zeeberg, B., Buetow, K.H., Schaefer, C.F., Bhat, N.K.,
Hopkins, R.F., Jordan, H., Moore, T., Max, S.I., Wang, J., Hsieh, F.,
Diatchenko, L., Marusina, K., Farmer, A.A., Rubin, G.M., Hong, L.,
Stapleton, M., Soares, M.B., Bonaldo, M.F., Casavant, T.L.,
Scheetz, T.E., Brownstein, M.J., Uedlin, T.B., Toshiyuki, S.,

TITLE
JOURNAL
PUBMED
2 (bases 1 to 4087)
AUTHORS
TITLE
JOURNAL

REMARK
COMMENT
NHL-MGC Project URL: <http://mgc.nci.nih.gov>
Contact: MGC help desk
Email: cgabs-remail.nih.gov
Tissue Procurement: ATCC
cDNA Library Preparation: Life Technologies, Inc.
DNA Sequencing by: Sequencing Group at the Stanford Human Genome
Center, Stanford University School of Medicine, Stanford, CA 94305
Web site: <http://www.bhg.stanford.edu>
Contact: (Dickson, Mark) mcd@paxil.stanford.edu
Dickson, M., Schmutz, J., Grimwood, J., Rodriguez, A., and Myers,
R. M.

Clone distribution: MGC clone distribution information can be found
through the I.M.A.G.E. Consortium/HLN at: <http://image.llnl.gov>
Series: IRAC Plate: 22 Row: m Column: 14
This clone was selected for full length sequencing because it
passed the following selection criteria: Hexamer frequency ORF
analysis
This clone has the following problem: retained intron.
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ORIGIN

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Matches 859; Conservative 1; Mismatches 544; Indels 69; Gaps 12;

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Dp	2357	-----TTTTTTTTTTTGAATAGAGCTGCTCTGTGCTGACGGCTGGAGT	2402
QY	1014	ACCTCAACCTCCGTTCACTGCTCACTGCAACCTCTGCTCCCGGGTTCAAGYAAATCTCCGTG	1073
Dp	2403	GCAGTGTGTGATCTCGGCTGCTGCAACCCCGCTCTCCGGGTTCAAGTATTTTCTG	2462
QY	1074	CTTAAGCCTCTGAGTAGCTGAAATTAACAACGTGACACACATGCTTGAGCTAAATTTTTT	1133
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QY	1134	GTAATTTTAGAGAGATGGGGTTTTACATGTGCCAGGCTGGTTCAACTCCTGAC	1193
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QY	1477	AACCTTAATGTACTGTGTTTTTCATGTGTACTAATTTTATGCTCACTGAATATATGCTTAA	1536
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Dp	2997	AATATTAATTTTGTGTACAAAGATTAATTTGTGTATCTATTAAGAAATACTTCAA	3056
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QY	1896	TACTCGGGAGGCTGAGGCAAGAAATTCCTTGAACCTCGGGAGGCGAGAGCTTGACGTGACG	1955
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Db	3353	CATGGTTGTGCACTGCACCTTCAGCGCTGGGGTGAAGAAGTAAAGCCCTGTCTAAAAAGCAA	3412
QY	2016	AAATTAATTAATTAATTAAGATCGGAGAGAAACA	2048
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HSM803026			
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DEFINITION	Homo sapiens mRNA; CDNA DKFPZ76100217 (from clone DKFPZ76100217).		
ACCESSION	AL713681		
VERSION	AL713681.1	GI:19584382	
KEYWORDS	HTC.		
SOURCE	Homo sapiens (human)		
ORGANISM	Homo sapiens		
	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Homidae; Homo.		
REFERENCE	1 (bases 1 to 3990)		
AUTHORS	Ansoerge,W., Krieger,S., Regiert,T., Rittmeller,C., Schwager,B., Wewes,H.W., Well,B., Amid,C., Obanger,A., Fobo,G., Han,W. and Wiemann,S.		
CONSRMT	The German CDNA Consortium		
TITLE	Direct Submission		
JOURNAL	Submitted (22-SEP-2004) MIPS, Ingolstaedter Landstr.1, D-85764 Neuherberg, Germany		
	Clone from S. Wiemann, Molecular Genome Analysis, German Cancer Research Center (DKFZ); Email s.wiemann@dkfz-heidelberg.de; sequenced by EMBL (European Molecular Biology Laboratories, Heidelberg/Germany) within the cDNA sequencing consortium of the German Genome Project.		
	This clone (DKFPZ76100217) is available at the RZPD Deutsches Ressourcenzentrum fuer Genomforschung GmbH in Berlin, Germany. Please contact RZPD for ordering: http://www.rzpd.de/cgi-bin/products/cl.cgi?cloneID=DKFPZ76100217 Further information about the clone and the sequencing project is available at http://mips.gsf.de/projects/cdna/.		
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CDS			
ORIGIN			

Query Match 13.6%; Score 341; DB 4; Length 3990;
Best Local Similarity 57.5%; Pred. No. 1.3e-21;
Matches 856; Conservative 1; Mismatches 541; Indels 91; Gaps 10;

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RESULT 4

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LOCUS 260L13-C56 CITB Homo sapiens genomic clone 260L13, genomic survey
DEFINITION
SEQUENCE.
A0839852
A0839852.1 GI:6652484

VERSION
KEYWORDS
SOURCE
ORGANISM

Homo sapiens (human)
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Hominidae; Homo.

REFERENCE
AUTHORS
1 (bases 1 to 15970)
Carpen, J.D., Nakalowska, I., Robbins, C.M., Scott, N., Sood, R.,
Comoros, T.D., Bonner, T.I., Smith, J.R., Faruque, M.U., Stephan, D.A.,
Pinkert, H., Morgenbesser, S.D., Su, K., Graham, C., Gregory, S.G.,
Williams, H., McDonald, L., Baxevanis, A.D., Klingler, K.W. and
Landes, G.M.

TITLE
A 6-Mb high-resolution physical and transcription map encompassing
the hereditary prostate cancer 1 (HPC1) region
JOURNAL
PUBMED
10708513
Contact: Carpen JD
Cancer Genetics Branch
National Human Genome Research Institute/National Institutes of
Health
Bldg. 36, Room 3D04, 36 Convent Drive, Bethesda, MD
Tel: 301 435 5626
Fax: 301 435 5465
Email: jdc@hgrl.nih.gov
Class: Shotgun.

COMMENT

FEATURES
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Location/Qualifiers
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ORIGIN

Query Match 13.5%; Score 337.2; DB 9; Length 15970;
 Best Local Similarity 57.8%; Pred. No. 1.1e-21;
 Matches 905; Conservative 1; Mismatches 529; Indels 131; Gaps 12;

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 Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
 Homidae; Homo.
 REFERENCE 1 (bases 1 to 897)
 AUTHORS NIH-MGC <http://mgc.nci.nih.gov/>.
 TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
 JOURNAL Unpublished (1999)
 COMMENT Contact: Daniela S. Gerhard, Ph.D.
 Office of Cancer Genomics
 National Cancer Institute / NIH
 Bldg. 31 Rm10A07 Bethesda, MD 20892
 Email: cgabs-remail.nih.gov
 Tissue Procurement: Dr. Michael Brownstein
 cDNA Library Preparation: Invitrogen Corp
 DNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
 DNA Sequencing by: Agencourt Bioscience Corporation
 Clone distribution: MGC clone distribution information can be
 found through the I.M.A.G.E. Consortium/LNL at:
<http://image.jnl.gov>
 Plate: ND4464 row: k column: 21
 High quality sequence stop: 636.
 Location/Qualifiers
 1. .897

FEATURES
 source

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/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone_image="303090116"
/lab_host="DH10B-Ton A ( T1 and T5 phage resistances )"
/clone_lib="NIH MGC 180"
/notes="Organ: Testis; Vector: pCMV-SPORT6.1; Site 1: NotI
site 2: EcoRV (destroyed) ; Library is oligo-dT primed and
directionally cloned (EcoRV site is destroyed upon
cloning). Average insert size 1.68 kb. Library was
constructed by (Invitrogen). Note: this is a NIH MGC
library."

```

[illegible]

KEYWORDS	HTC.
SOURCE	Pongo pygmaeus (orangutan)
ORGANISM	Pongo pygmaeus Eukaryote; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eultheria; Euarchontoglires; Primates; Catarrhini; Hominoidea; Pongo.
REFERENCE	1 (bases 1 to 3095)
AUTHORS	Bloecher,H., Boecker,M., Brandt,P., Mewes,H.W., Well,B., Amid,C., Osanger,A., Fobo,G., Han,M. and Wiemann,S.
CONSRFM	The German cDNA Consortium
TITLE	Direct Submission
JOURNAL	Submitted (12-NOV-2004) MIPS, Ingolstaedter Landstr.1, D-85764 Neuherberg, GERMANY
COMMENT	Clone from S. Wiemann, Molecular Genome Analysis, German Cancer Research Center (DKFZ); Email s.wiemann@dkfz-heidelberg.de; sequenced by GBR (National Research Centre for Biotechnology Ltd., Braunschweig/Germany) within the cDNA sequencing consortium of the German Genome Project.
FEATURES	This clone (DKFZp469P012) is available at the RZPD Deutsches Resourcenzentrum fuer Genomforschung GmbH in Berlin, Germany. Please contact RZPD for ordering: http://www.rzpd.de/cgi-bin/products/cl.cgi?cloneID=DKFZp469P012 Further information about the clone and the sequencing project is available at http://mips.gsf.de/projects/cdna/.
SOURCE	Location/Qualifiers 1..3095

Query Match	12.4%	Score 311;	DB 4;	Length 3095;
Best Local Similarity	56.6%	Pred. No. 6.1e-19;		
Matches	883;	Conservative 1;	Mismatches 511;	Indels 166; Gaps 10;
QY	584	GTAAAGTGTGACAGTTGGGTTTTTTTTTTTGTGTGTGTTTAAAGACAGGGTCTTGCTCTGT		643
DB	1570	GTGAGCCACTACCTGTGACGCTTTTGTGTGTGTTTAAAGATGAGTCTCACTTGT		1629
QY	644	CACCAAGGATGAGACACAGTGTGTCAACCATAGGTCACTGACAGCTCAACCTCTGAGCT		703
DB	1630	TGCCCACTTGAGATGACAGTGTGTGCACTTCACTCACTGACAGCTCCGCTCCCAAGTT		1689
QY	704	CAAGGATCTGTGACTCTGAGCTCTCCCAAGTAGTGTGGAGCTACGAGCTGACCAACAG		763
DB	1690	CAAGCAATTCCTGCGCTCAAGCTCTCCCAAGTAGTGTGGAGCTGTGTGACCAACCA		1749
QY	764	CTGTGCTAATTAATAAATTTTTTTTGTAGAGACTGGGCTTACTACGTTGGCAAGGCTTG		823
DB	1750	CCAGCTAATTTT---TTTGTATTTTATGGAAGCAGGGTTTCAACATGTGCTCCAGGCTGG		1806
QY	824	TCTTAACTCTGTGGCTTCAGCAATCTCTACCTTGGCATCCAAAGTGTGGAGTTAC		883
DB	1807	TCTTGAATCTGTGAGCTCAGGACATCCGCGCCACCTCAGCTCTCCCAAGGTGCTAGGATTAC		1866
QY	884	AGGGGTAGCCACCATGTGGCGGCTACTATTCTTTACATTC-----		925

Db	1867	AGGCTAAGCCACCATGCCCTCGCATGTAAGCCCTTTTGGACATTACAGATATTCGCTA	1926
Oy	926	-----CATCTTCCAT-----A	938
Db	1927	TTTTCGAGGGAGCTCAAGTAATGCTATAGACGAACTTTCGAATATCTCAATGTTGA	1986
Oy	939	GAATGTAGATCCACAGAACAGGATATCTGCTATTTTCTCCCTTCTTTTGGACA	998
Db	1987	AATATTGGAGAACAAACTTGTTTACATACCTTTTTTTTTTTTTTTTTTTTTTGGAGCC	2046
Oy	999	GAGT-----CTCATCTTCAACCTCAACCTCCGGTCACTCACTGCACAC	1043
Db	2047	GAGTTTGTCTTGTGTCGCCAGATCTGGAGTGAATGCGCCCAATCTCACTCACTGCACAC	2106
Oy	1044	TCCTGCTCCGGGTTCAAGYAATCTCTGCTTAAGCCTCTGTAGTACCTGAATTACAA	1103
Db	2107	TCCTGCTCCAGGTTCAAGCAATCTCTGTCTGAGCTCCCGATGACTGGATTACAG	2166
Oy	1104	GCGTGCACCACTTACTGCTGCTAATTTTGTATTTTTAGACAGATGGGTTTACAT	1163
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Oy	1164	GTGCGCCAGCGTGTCTAACTCCGTACTCAAGTATCTGCGCTGCTCACTGCTCCCA	1223
Db	2226	ATTGTCTACGGCTGTTCTAACTCTGACCTTCAGACGATCGGCTGCTCAACTTCCA	2285
Oy	1224	AGTGTGAAATTATAGCGTGAAGTCACTGCTGCGCC-GATTACTGTCTATTTTCTTTA	1282
Db	2286	GTTGCTGGGATTAATGCTGAGCCACACACCGGCTGCTTAACTATATCTTGACA	2345
Oy	1283	TTGCTATATCCCAAGATTAAGACATGTCTGAATATATAGTGTCTCAATTAATAT	1342
Db	2346	TTTCTAGCTGAAGATTATAGGTGATGTTGGGAGTGGAGTCAATGAATAAT	2405
Oy	1343	GATGAATGCAAGCCTGATATTAACCTTCTTTTCTTTTAAACAATCTTGACAC	1402
Db	2406	TGAGTACAGACATGCCAGAGACA-----TTGGGGGCGAGATGATCGGCACT	2454
Oy	1403	TTTGCAGATTAATATACATCTTGATTCGCTTTTCACTTATCACTTGTATGACTTT	1462
Db	2455	GATGTGATGGAATGGCAGTGGAGGTGGAGGAACAGATATCAATCCAGATGGAAGT	2514
Oy	1463	TTCAATTTGCTCAAACTTATTTGTAATGTTTTTCAATGTTACTATTTTACTCTG	1522
Db	2515	AGAACCCCTGACTCAAGTCACTGTGGGAATGAAAAAGAGATTATTTGAGACTA	2574
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Db	2575	CCATTAAGTCAAGAAATGTTTGAATGGACAGAAATGACTTAAGAGGGAGTCAATGG	2634
Oy	1583	AAATCTGATGAAGCTATGAACCTCTCCAGAGAAATACACACACACACACTCA	1642
Db	2635	TA-----ACCC	2641
Oy	1643	CACACAGTTTTTTTAAATGTTGCAACTAAGACAGAAACCTGCAATTAAGATGTTTG	1702
Db	2642	AAGCAGAGGTTTTCAGATTTTCTCAGGTAGAAGCTCCCATAGAGAGATTG-----	2691
Oy	1703	TTCAATATTAATTAATACTCACTGATGGGACAGTGACTTAACCTGTATCCACACTAC	1762
Db	2692	-----TTTAAATGAAGCTTCTGGGCTGGGTGTGGCTCAACCTGTATATCCACAG	2745
Oy	1763	TTTGAAGTCCAAAGTGGTGGATCACTTGAGTGAAGATTTGAGACAGGCTGCTCAA	1822
Db	2746	TTTGGAGGCGATGGCAGGCGGATTTGGGTAGAGTCAAGATTTCAAGACAGCTTGGGAG	2805
Oy	1823	TATGTGAACCCATCTCTACTAATAAATACAAAATTAAGCTGGGTATGATGATGC	1882
Db	2806	CATGTGAAACCGTATCTC-ACATAAATAAATAATTAAGTTGGGTGTGGGCAAGCAC	2864
Oy	1883	CTGTAGTCCCACTCTCGGAGGCTGAGGACAGAAATGCTTGAACCTGGGAGGCGA	1942

Df		2865	CTGTAATCCCAAGCCTCTTAGGCACGTAGAGAGAAGAATCATCTTGAACCTGGAGGCGA	29224
Oy		1943	GATTGCACTGAGCCGAGATCCACCACCTGCACTTCACGCTGGGCGACAAGCGACTCT	20020
Df		2925	GGTTGACAGTGAGCCAAAGTACGCCACCTGCACTCACGCTGGGCGAACAAAGTAAACTCT	29844
Oy		2003	ATCTCAAATAAATTAATTAATTAATTAAGATGCGAGAGAAACAAATTAAGATTC	20623
Df		2985	GTTCTCAAAAAAAAAAAAAAAAAAAAAAACAAGCTAAGTATCAAGCTCTTC	30444
Oy		2063	C 2063	
Df		3045	C 3045	
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LOCUS	CR749233/c	5797 bp	mRNA	linear
DEFINITION	Homo sapiens mRNA; cDNA DKFPZ686C0331 (from clone DKFPZ686C0331).			
ACCESSION	CR749233			
VERSION	CR749233.1			
KEYWORDS	HTC.			
SOURCE	Homo sapiens (human)			
ORGANISM	Homo sapiens			
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Homnidae; Homo. 1 (bases 1 to 5797) Poustka,A., Albert,R., Moosmayer,P., Schupp,I., Wellenreuther,R., Mewes,H.W., Weil,B., Amid,C., Osanger,A., Fobo,G., Han,M. and Wiemann,S. The German cDNA Consortium Direct Submission Submitted (17-AUG-2004) MIPS, Ingolstaedter Landstr.1, D-85764 Neuerberg, GERMANY			
COMMENT	CONSRMT TITLE JOURNAL			
FEATURES				
source				
gene				
CDS				
locus_tag				
cds_start				
cds_end				
cds_frame				
protein_id				
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		/tissue_type="heart"
		/clone_lib="468 (synonym: phrcl). Vector pSport1_sfi; host DHIOB; sites SfilA + SfilB"
		/dev_stage="adult"
		/note="unclassified"
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Dd	1602	TTTTTTTTTTTTTTTGTTCCTTTGTTTGAAGACGAATCTTGCTCTGTCTCCCGGCTGAG 1543
Oy	658	CACAAGTGTGCAACCATAGTCACTGACGACTCAAACCTCTAGACTCAGAAGAACTGCTG 717
Dd	1542	TGTATGTGTGAATCTCAGCTCACTGCAAACTCATCTCCGGATTCACTTAATCTCTCT 1483
Oy	718	ACCTGAGCTTCCCAGTAGTACTGCGGACTAGACGCTGACCCAACAGCTGCTTAATAAA 777
Dd	1482	GCCCTAGCTCTCAGTAGTACTGCGGACTATGAGGCTGCTGCCATAATGCCCCAACATAATTT 1423
Oy	778	AAAATTTTTTTGTAGAGCTGGGCTTTACTAGTGGGCTGAGGCTTGTCTTAACCTCCCTGG 837
Dd	1422	GTA--CTTTATGATGAGAGCGGGGTTTACACAGTGTGGCAGGCTGAGTCTAACTCTCTGG 1365
Oy	838	CTTCAAGCAATCTCTCTCACTCTGTGCATCCCAAAGTCTGCGGATTACAGGGGTGAGCCACC 897
Dd	1364	CCTCAAGTAGATCCACCTGCTGCTTGGCTCTCCAAAGTCTGGGATTACAGGTGTGACCACTG 1305
Oy	898	ATGTGCGGCTACTATTTCTTTAATTCCATCTTTCCAAATGAAATGTAAGATCCACGAA 957
Dd	1304	CACCTGGCTGAATTTCTCCAAATCTTCCCA-----CACACCCGCTCAGTCTCTCTTCC 1250
Oy	958	CAGGATATACGCGCATATTTCTCTCTTTTGTGAGACAGAGTTCACATTCACTCACT 1017
Dd	1249	TGATCATTTAGACGCTTTTTTTTTTTTTTTTTTTTGTGAGACAGCAATTCACATCTCACTCC 1190
Oy	1018	CAACCTCCGCTCA-----GCTCACTGCAACCTCTGCTCTCCGGGTTCAAGY 1063
Dd	1189	AGACTGAGTGCAGAGTAGTCAATCTTGCTCACTGAACCTCTTCTCTCCAGCTCANAGC 1130
Oy	1064	GATTCTCTGCTTAAGCCTCTCTAGTAGCTGAAATTACAGACGTCACCAACCATGCTTGG 1123
Dd	1129	GATTCTCTGCTCAAGCCTCTCCAGTAGCTGGGATTACATGTGTGACCACTAACACCA 1070
Oy	1124	CTAATTTTTTGTATTTTGAAGAGATGGGTTTACATGTGTGCCAGGCTGCTCTCA 1183
Dd	1069	GATAATTTTTTGTACTTTTAGTAGAGATGGGTTTGAACATATTGGCCAGACTGTGCTTGA 1010
Oy	1184	ACTCTGACTCTCAAGTAGTATGCTGCGCTCAAGTCTCCCAAAGTGTGGAAATTATAGGCGT 1243
Dd	1009	ATTCTTGACTTCAAGTAGATCCGCCCACTTCAAGCTCTCCAAAGTGTGGGTTATAGGCGT 950
Oy	1244	GAGTCACTGTGCTGCGCATTAATCACTATCTTAATTTCTTTAT 1283
Dd	949	GAGGCACTGCAACCGGCCCCCTCTCTTCATCTTAGTCAAT 910
RESULT 9		
CR59842	1663 bp	mRNA linear HTC 21-JUL-2004
LOCUS	full-length cDNA clone CSDBA002Y006 of Neuroblastoma of Homo sapiens (human).	
DEFINITION	CR59842	
ACCESSION	CR59842.1	GI:50480649
VERSION		

	KEYWORDS	HTC; CNSLT cDNA.
SOURCE	Homo sapiens (human)	
ORGANISM	Homo sapiens	
	Eukaryota; Metazoa; Chordata; Craniata; Vertebrate; Euteleostomi;	
	Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;	
	Hominidae; Homo.	
REFERENCE	1 (bases 1 to 1863)	
AUTHORS	L.I. W.B., Gruber, C., Jesssee, J. and Polayes, D.	
TITLE	Full-length cDNA libraries and normalization	
JOURNAL	Unpublished	
REMARK	Contact : Feng Liang Email : fliang@lifetech.com URL : http://fulllength.invitrogen.com/Invitrogen Corporation 1600 Faraday Avenue 2 (bases 1 to 1863) Genoscope. Submitted Submission Submitted (20-JUL-2004) Genoscope - Centre National de Sequencage ; BP 191 91006 EVRY cedex - FRANCE (E-mail : sequef@genoscope.cns.fr - Web : www.genoscope.cns.fr) 1st strand cDNA was primed with a NotI-oligo(dt) primer. Five primers into the Not I and EcoR V sites of the pCMVSPORT 6 vector. Library was normalized. Library was constructed by Life Technologies, a division of Invitrogen.	
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Best Local Similarity	56.9%; Pred. No. 6e-18;	
Matches	811; Conservative 1; Mismatches 509; Indels 105; Gaps 10	
Db	624 GAGACAGGGTTCTCTGTCAACCAGGCATGAGCACAGTGTGCACCATAGTCACGTG 683	
	496 GGGTCAGAGTCTCACTGTTCCTCCAGGCTGGAAGTCTGTGGCACCGTGTGGCTACTG 555	
Oy	684 CAGCCTCAACTCTGAGCTTAAGGATCTGTGACTCAAGCTTCCAAGTAGCTGGAC 743	
Db	556 CAACCTCGTCTCTGGGTTCMAACAATTCTTGCCCTCAAGTAGCTGGAGAT 615	
Oy	744 TAGAGGCTGACACCAACGCCCTGGCTAATTAATAAAATTTTTGTAGAGCTGGGCT 803	
Db	616 TACAGGTCCACCAACCAACCCGCTAATTTTTTATATTTTATATGACACGGGGTTT 675	
Oy	804 TACTACGTTGGCCAGGCTTGTCTTAAATCCTGGCTTCAGAACAAATCCTCACTTGACA 863	
Db	676 CACCATGTGGCCAGGCCGCTTGAATTGTGAAGTAGAGTAGACACCGGCTTGGCC 735	
Oy	864 TCCCAAAGTGTGGATTAACGGGGTGAACCAACATGTGGGCTATCTTATTTCTTAAT 923	
Db	736 TCCCAAAGTGTGGGATTAACAGTTCAGATTGTAACAAATCTGCTAGAGATTGAACAG 795	
Oy	924 TCCATCTTTCCAAATGAATGTAAATCCACAGAACAGGAGTTATCTGCCTATTTTCTT 983	
Db	796 AGCTGCCTGGAAATCCCGCCACAGTGAAGAGACCTGAAGAACAAGAAAACAACGAAC 855	
Oy	984 TTCTTTTGTGAGACAGAGTCTCACTTCATCACTCAACCTCCGTTCAAGTCACTGCAAC-C 1044	
Db	856 GGAATCTTGTCTGTGTGTCAAGCTGAGAGTGAATGTGGCAATCTCGGCTCCCTGCAAGA 915	
Oy	1043 CTGTGCTCCCGGATTCAGATTTCTCTGCTTAAGCTTCTCTGAAGTAGCTGAATTACA 1100	
Db	916 ATGTGCTCCCAAGTTCACGCCATTTTCCCACTCAAGCCCTGAGAGTAGCTGAAGTACA 975	
Oy	1103 AGCGTGACCAACCATGCTGGCTAAT----TTTGTATTTTGTAGCAGATGGGGTTT 1156	
Db	976 GGCTTCGGCCACCAACGCTGGCTAATTTTGTGTGTATTTTGTATGAGACGGGGTTTC 1038	

QY	1159	ACCAATGTTGCCAAGCTGCTGCTCAAACTCCGACCTCAAGGAACTCTGCGCCCAAGCT	1218
Db	1036	ACTGTGTAGGTAGAGATGCTCAATCTCCGACCTCA--TGATTCATCCGCTTGCGCT	1093
QY	1219	CCCAAGTGCAGGAATTATAGCGGTGATCACTGCTGCGCCGATTAAGTCTCATTTTC	1278
Db	1094	CTCAAAAGTCTGGGATTACAGGCATGAGCAACCGAAGCTGGCTG-----TTTT	1141
QY	1279	TTTATTTGCTATATCCCAAGATCTAGAGAGTGTCTGACATATAGTAGTGTCTCAATAAT	1338
Db	1142	TTCAATTTCTAATTATGATTTGTGTCTGGAAATGACAGTACTTAAGGGGTTCATCATG	1201
QY	1339	AATGATGAATGACACAGCCCTAGATATAAATCTTCTTTTCTTTTTPAAAACAATCTGA	1398
Db	1202	TTTATTAAGCTGAATAGTGTCTGTATGTGACGTACAGACACCCTAGTGTACATGCTACT	1261
QY	1399	CAACTTGGCAGAAATAAATACAACTTGCATTTCTGTTTTCATTATCACTTGTTATGA	1458
Db	1262	CTCCGTGCTGCATAGATGGGGGCGCTGCCACACCAAAATCTCCGGGTTCGCCGTGCC	1321
QY	1459	CTTTTTCATATGCGTCAAAACCTTTATTTGTATAGTGTTTTTCATTTGTAATTTATGTC	1518
Db	1322	GCACCTGAAGGTACCCACCAAGTGTCTGTCACTGCGCTCAATCCACTTTCTGTGTGT	1381
QY	1519	ACTGAATTAATATGCTTAATTTGCTTATACATCTCTCGCTCACTTTAGAAGGCCAAT	1578
Db	1382	TCTCATGTTTCTGCGCTC-----CCTAGAGACATTTCTCTCTACAGAGTAG-----	1428
QY	1579	TTACAAATCTGATGAAGCTATGAACCTCTCCGAGAAATACACACACACACACACA	1638
Db	1429	-----TTTCAATGTAAATACATATGCCAGACACATTA	1459
QY	1639	CTCACACACAGTTTATTTTATGTTTGCACCTAAGACAAAGAACTGECATTAGAGATG	1698
Db	1460	C-----GTCAATTTAGATTTTCTAGTAGTCACTTAGAAAAGTAAAAAAGGGCCA	1510
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Db	1511	GCTG-----CAGTGACTCACACCTGTATATCCA	1538
QY	1759	GTACTTTGGAAAGTCCAAAGGTGGGTGGATCATCTGGAGTGAAGTTGCAGACGAGCTGG	1818
Db	1539	GCACTTTGAAAGGCTGAGGTGGTGGATCAC--GAGGTCAAGAGATCGAAGACATCTGG	1566
QY	1819	TCAATATGSGTGAACCCCTATCTCTACTATAAAATAC-AAAAATTAAGCTGGGTGTATGATG	1877
Db	1597	CCAACATGSGTGAACCCCTCTCTACTATAAAATACAAAAATTTAGCTGGGCGTGGTGCA	1656
QY	1878	CATGCGCTTAATGCCAGCTACCTCGGGAAGCTGAAGCAGAGAAATTGTCTTGAACCTGGAG	1937
Db	1657	GAGCGCTGTAGTCCAGCTACTCGAGAGGCTGAGGCAAGAGAAATGGCTGAAACCGGTAG	1716
QY	1938	GCAGAGTGTGCAGTGAGCGGAGATCCACCACTGCATCTCAGCCTGGCGCACACGCGAG	1997
Db	1717	GCAAGAGCTTGAGGTAGCGAGCATAGCGCATCTGCATCTCAGCCTGGCGCACAGGGAG	1776
QY	1998	ACTCTATCTCAAAAAATTAATTAATAAATTAAGATCGGAGAGA	2043
Db	1777	ACTCATCTCAAAAAAAGGTTTACAAAGAAACAGGTGAAA	1822

RESULT	10
BQ958903	
LOCUS	BQ958903 946 bp mRNA linear EST 21-AUG-2002
DEFINITION	AGENCOURT_100354B5 NIH_MGC_40 Homo sapiens cDNA IMAGE:6483214 5' mRNA sequence.

ACCESSION	BQ958903	
VERSION	BQ958903.1	GI:22374381
KEYWORDS	EST.	
SOURCE	Homo sapiens (human)	
ORGANISM	Homo sapiens	
	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;	

REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT

Mammalia: Eumetaria: Euarchontoglires; Primates; Catarrhini;
Homidae: Homo.
1 (bases 1 to 946)
NIH-MGC <http://mgc.ncbi.nlm.nih.gov/>.
National Institutes of Health, Mammalian Gene Collection (MGSC)
Unpublished. (1999)
Contact: Robert Strausberg, Ph.D.

Email: cgabbs-remail.nih.gov
Tissue Procurement: DCTD/DTF
cDNA Library Preparation: Rubin Laboratory
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
DNA Sequencing by: Agencourt Bioscience Corporation
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LNL at:
<http://image.llnl.gov>
plate: L10C667 row: g column: 23
High quality sequence stop: 534.

FEATURES
source

Location/Qualifiers
1. .946

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organism="Homo sapiens"
mol_type="rRNA"
/db_xref="taxon:9606"
/clone="IMAGE:6483214"
/tissue_type="carcinoma, cell line"
/lab_host="DH10B (phage-resistant)"
/clone_lib="NIH MGC 40"
/notes="Organ: prostate; Vector: pOTB7; Site_1: XhoI; Site_2: EcoRI; cDNA made by oligo-dT priming. Directionally cloned into EcoRI/XhoI sites using the following 5' adaptor: GGCAACAG(G). Library constructed by Ling Hong in the laboratory of Gerald M. Rubin (University of California, Berkeley) using ZAP-cDNA synthesis kit (Stratagene) and Superscript II RT (Life Technologies)."
Note: this is a NIH MGC library."

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ORIGIN

Query Match	12.0%;	Score 300.6;	DB 5;	Length 946;
Best Local Similarly	69.1%;	Pred. No. 1e-17;		
Matches 468, Conservative	1;	Mismatches 197;	Indels 11;	Gaps 4;

596 AGTTTGGGTTTTTTTGTTCCTTTAGAGACAGGGCTTTGCTCTGTCCACCCAGGCATG 655

D_b

24 AGNNCTTTTGTGTTTTGGAGACAGAATCGTCCTGCCTCATCCCAAGCCTGG 83

[illegible]

656 AGCACAGTGGTGGACCAATGGTACATGACGCTTCATCCCTCTGGATCATGGGATCTGC /13

Db 84 AGTGCAGTGGTG---CGATCGCTCACTGCACACTCAGCCTCCCTGAGTTAAGTGATTCTC 140

716 TGACCTAGCCTCCCAAGTAGCTGGACTACGAGCGTGACCAACCAAGCCTGGCTAATTA 775

Db 141 CTGCCTCAGCCTCCCAAGTAGCTGGGATTACAGGTGTGCACCACCATGCCCCGGTAATTT 200

776 AAAAAATTTTGTAGAGCTGGGCTTTACTACGTTGGCCAGGCTTGTCTTAAACTTCCT 835

Db 201 TTTTGTATTTTAACTGAGACAGGCTTACCATGTTGGCCAGGCTGCTCGAAGTCTT 260

836 GAGTTTAAAGCAATCTCTCCATCTTGCCATCCCAAGTGTCTGGATTACAGGGGTGACCA 895

[illegible]

DB
Z6L GARCIC--G7WALCCGCCICCCACBCCCCCCHHAGVGCIGGGHILPCACBAGVIGIWBURCN ZIO

896 CCATGTGCGGCTACTATTCTTTACATTCCATCTTTCCATAGAGTAGATCCACAG 955

Db 319 CCACGCGTGGCCCCAGTCCAAATATTTAAAGATTGTTCCTTAGTGTCTTGAAGCTTTTC 378

956 AACAGGATTA CTG CCTATTTCTTCTTTTGGAGACAGCTCACTCATCAC 1015

Db 379 CACA-----AAATCTTTTTTGAGATGAGTCTCACTCTGTCAACCAGGCTGAGTGC 433

1016 CTCACCTCCGTTACGCTACTGCACCTCTGCTCCCGGGTCAAGYGATTCTCTGCC 1075

[illegible][illegible]

1076 TAAGCTCTGAGTAGCTGGMTACAAAGCGTCACCACTCTGGCTAA1111161 1133

Plate: NDAMI172 row: C column: 08
 Seq primer: M13-21 (TGTAAACGACGGCCAGT)
 High quality sequence stop: 711
 POLYA=yes.

FEATURES

Source

Location/Qualifiers

```

1..736
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:30928759"
/sex="male"
/class_type="human embryonic stem cells"
/cell_type="human embryonic stem cells"
/lab_host="DH10B-T1 phage-resistant E. coli"
/clone_lib="NIH MGC 260"
/notes="Vector: pExpress-1; Site 1: NotI; Site 2: EcoRV;
RNA obtained from human embryonic stem cells isolated from
the inner cell mass of blastocyst stage embryos. Cell line
id and NIH Registry designation is BG01. Positive for
SSEA3, SSEA4, Tra 1-60, Tra 1-81, CD9, Alk Phos, Oct4 and
Nanog expression; negative for SSEA1 expression. Passage
number 21. cDNA primed using oligo-dT primer:
5'-pGACTAGTTCTAGATCGGAGCGGCCCT(T)25-3 and cloned into
the EcoRV/NotI sites of pExpress-1. This primary library
is non-normalized (normalized primary library is
NIH MGC 261). It was constructed by Express Genomics
(Fredrick, MD). Sequence ends have been trimmed to
exclude vector and regions below phred quality 16. Note:
this is a Mammalian Gene Collection library."

```

ORIGIN

Query Match 11.7%; Score 293.6; DB 8; Length 736;

Best Local Similarity 66.9%; Pred. No. 5e-17;

Matches 464; Conservative 1; Mismatches 220; Indels 9; Gaps 3;

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QY 599 TTGGGTTTTTTTGTGTTTGAAGACAGGGTCTGCTGTGACCCAGGATGAGC 658
Db 1 TTTTGTGTTTGTGTTTGTGTTTGAAGATGAGTCTCACTCTCTGCGCCAGGCTGAGT 60
QY 659 ACAGTGTGCAACCTAGTCACTGCAACCTTCACTTCAAGCTCAAGGATCTGCTGA 718
Db 61 GCAGTGGGGGATCGGCTCACTGCAACCTTCTGCTTGGTTCAAGCACTTCTCTG 120
QY 719 CCTGAGCTCCCAAGTACTGAGGACTAGAGGCTGACACCAAGCTGCTAATTAATA 778
Db 121 CTTAGAGCCCTGAGTACTGGATTAACAGCACCAACCAAGCTGAGTACTT--TT 178
QY 779 AAATTTTGTGAGAGCTGGTCTTACTAGCTTGGCAGGCTTGTCTTAAACTCTGGC 838
Db 179 TTGTATTTTGTGAAAAATGGGGTTTGCTATATTGGCAGGCTGGTCTCAAACTCTGAC 238
QY 839 TTCAAGCAATCTCTACTTGGGATCCCAAGTCTGGGATTAAGGGGTGAGCCACA 898
Db 239 CTCAGGTATCACTGCTGAAAGCTGCAAGTCTGGGATTAAGGACATGAAACACA 298
QY 899 TGTGGGCTACTTATTTCTTACATTCATCTTCCATTAAGATTAAGATCAAGAAC 958
Db 299 TGCCAGGCTCTTATTTCTTTTAATTAATATGACAAGTTATTTATTTATTTA 358
QY 959 AGGGAATACGCTATTTCTCTCTTTTGTGAGACAGTCTCACTTCATGACCTC 1018
Db 359 TTTATTTATTTTGAAGATGAGTTTGCCCTTGTTGCCCAAGCTGAGTGAAATGGC 418
QY 1019 AACCTCCGTTGAGCTCACTGCAACTCTGCTCCGGGTTCAAGATTTCTCTGCTCA 1078
Db 419 GATC-----TCAGCTCACTGCAACTCACTCCGGGTTCAAGATTTCTCTGCTCA 473
QY 1079 GCTCTCTGAGTGTGATTAATTAAGAGCTGACCAAGCTTGGCTATTTTGTATTT 1138
Db 474 GCTCTCTGAGTGTGAGTGTGATTAAGAGCTGACCAAGCTTGGCTATTTTGTATTT 531
QY 1139 TTTAGCAGATGGGGTTTTTACATGTGGCCAGGCTGGTCTCAAACTCTGACCTCA 1198

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```

Db 532 TTTAGTAGAAGACAGAGTTTCTTCATGTGTCAGGCTGGTCTTGAACCTCTGAC 591
QY 1199 TGAATGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTG 1258
Db 592 TGAATGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTG 651

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```

QY 1259 GCCGATTACTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTG 1292
Db 652 GCCATTATCAAGATTATTTACTTCTCTCTC 685

```

RESULT 13

CR613629

LOCUS

DEFINITION full-length cDNA clone CS0D1039YN19 of Placenta Cot 25-normalized

of Homo sapiens (human).

ACCESSION

CR613629

VERSION

CR613629.1

KEYWORDS

HTC; CNSLT_cDNA.

SOURCE

Homo sapiens (human)

ORGANISM

Homo sapiens

REFERENCE

AUTHORS

TITLE

JOURNLT

REMARK

REFERENCE

AUTHORS

TITLE

JOURNLT

REMARK

REFERENCE

AUTHORS

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http://fulllength.invitrogen.com/InvitrogenCorporation1600

Faraday Avenue

Genoscope.

Submitted (20-JUL-2004) Genoscope - Centre National de Sequencage :

BP 191 91006 EVRY cedex - FRANCE (E-mail : seqref@genoscope.cns.fr

- Web : www.genoscope.cns.fr)

1st strand cDNA was primed with a NotI-oligo(dT) primer. Five prime

end enriched, double-strand cDNA was digested with Not I and cloned

into the Not I and EcoR V sites of the pCMVSPORT 6 vector. Library

was normalized. Library was constructed by Life Technologies, a

division of Invitrogen.

Location/Qualifiers

1..3092

/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

/clone="CS0D1039YN19"

/class_type="Placenta Cot 25-normalized"

/plasmid="pCMVSPORT_6"

ORIGIN

Query Match 11.6%; Score 290.6; DB 4; Length 3092;

Best Local Similarity 69.7%; Pred. No. 3.7e-117;

Matches 466; Conservative 1; Mismatches 190; Indels 12; Gaps 5;

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QY 594 ACAGTTGGGTTTTTTTGTGTTTGAAGACAGGGTCTGCTGTGACCCAGGCA 653
Db 1750 AAATTTTGTGTTTGTGTTTGAAGATGAGTCTGCTGTGACCCAGGCT 1809
QY 654 TGACACAGTGTGCAACCATAGTCACTGACGCTCAACTCTGAGCTCAAGGATCT 713
Db 1810 GAGTGTGAGTGTG---CGATGCTCACTGCAACTCAAGCTCTGATTTAAATGATTC 1866
QY 714 GCTGACTGACCTCCCAAGTGTGGAAGTCAAGAGGTGACACACAGGCTGGTAAAT 773
Db 1867 TCTGCTGACCTCCCAAGTGTGGAAGTCAAGAGGTGACACACAGGCTGGTAAAT 1926
QY 774 TAAATAATTTTGTGAGAGCTGGGCTTACTAGTGTGAGGCTGCTTAAATCTC 833
Db 1927 TTTTGTGA-TTTTGTGAGAGAGAGGGTTTCAACATGTGGCCAGGCTGGTCTGMACTC 1985

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